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*      8228      8327: gap of 100 bp
*      8328      9072: contig of 745 bp in length
*      9073      9172: gap of 100 bp
*      9173      9920: contig of 748 bp in length
*      9921      10020: gap of 100 bp
*      10021      10740: contig of 720 bp in length
*      10741      10840: gap of 100 bp
*      10841      11566: contig of 726 bp in length
*      11567      12407: contig of 741 bp in length
*      12408      12507: gap of 100 bp
*      12508      13254: contig of 747 bp in length
*      13255      13354: gap of 100 bp
*      13355      14095: contig of 741 bp in length
*      14096      14195: gap of 100 bp
*      14196      14933: contig of 738 bp in length
*      14934      15033: gap of 100 bp
*      15034      15755: contig of 722 bp in length
*      15756      15855: gap of 100 bp
*      15856      16584: contig of 729 bp in length
*      16585      16684: gap of 100 bp
*      16685      17433: contig of 749 bp in length
*      17434      17533: gap of 100 bp
*      17534      18286: contig of 753 bp in length
*      18287      18386: gap of 100 bp
*      18387      19127: contig of 741 bp in length
*      19128      19227: gap of 100 bp
*      19228      19937: contig of 710 bp in length
*      19938      20037: gap of 100 bp
*      20038      20750: contig of 713 bp in length
*      20751      20850: gap of 100 bp
*      20851      21574: contig of 724 bp in length
*      21575      21674: gap of 100 bp
*      21674      22395: contig of 721 bp in length
*      22396      22495: gap of 100 bp
*      22496      23225: contig of 730 bp in length
*      23226      23325: gap of 100 bp
*      23326      24072: contig of 747 bp in length
*      24073      24172: gap of 100 bp
*      24173      24902: contig of 730 bp in length
*      24903      25002: gap of 100 bp
*      25003      25746: contig of 744 bp in length
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*      29877      29976: gap of 100 bp
*      29977      30717: contig of 741 bp in length
*      30718      30817: gap of 100 bp
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*      31555      31654: gap of 100 bp
*      31655      32380: contig of 726 bp in length
*      32381      32480: gap of 100 bp
*      32481      33227: contig of 747 bp in length
*      33228      33327: gap of 100 bp
*      33328      34072: contig of 745 bp in length
*      34073      34172: gap of 100 bp
*      34173      34896: contig of 724 bp in length
*      34897      34996: gap of 100 bp
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*      35740      35838: gap of 100 bp
*      35839      36559: contig of 720 bp in length
*      36560      36659: gap of 100 bp
*      36660      37395: contig of 736 bp in length
*      37396      37495: gap of 100 bp
*      37496      38105: contig of 610 bp in length
*      38106      38205: gap of 100 bp

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*      38206      38948: contig of 743 bp in length
*      38949      39048: gap of 100 bp
*      39049      39799: contig of 751 bp in length
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*      40647      40746: gap of 100 bp
*      40747      41467: contig of 721 bp in length
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*      41567      42307: contig of 740 bp in length
*      42308      42407: gap of 100 bp
*      42408      43130: contig of 723 bp in length
*      43131      43230: gap of 100 bp
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*      43954      44063: gap of 100 bp
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*      44772      44871: gap of 100 bp
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Query Match      100.0%; Score 200.6; DB 12; Length 194321;
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QY      61 CACACCAAGAGGAACTCGAATCTCATGTTGAGAAATGAAAGTGTAGAGCTGTTCT 120
DB      10510 CACACCAAGAGGAACTCGAATCTCATGTTGAGAAATGAAAGTGTAGAGCTGTTCT 10451
QY      121 TGGAGCCCAACAAGCAGATATTGCTTCTTAAGCTTAAGCAGATCTGCTACCTGTG 180
DB      10450 TGGAGCCCAACAAGCAGATATTGCTTCTTAAGCTTAAGCAGATCTGCTACCTGTG 10391
QY      181 GTCTTTCACCCCAAGCTGTGCA 201
DB      10390 GTCTTTCACCCCAAGCTGTGCA 10370

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RESULT 3
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DEFINITION Human plasminogen gene, exon 16.
ACCESSION M34272 J05286
VERSION M34272.1 GI:190060
KEYWORDS plasminogen.
SEGMENT 20 of 24
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

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REFERENCE 1 (bases 1 to 331)
AUTHORS Petersen,T.E., Mattzen,M.R., Ichinose,A. and Davie,E.W.
TITLE Characterization of the gene for human plasminogen, a key proenzyme in the fibrinolytic system
JOURNAL J. Biol. Chem. 265 (11), 6104-6111 (1990)
PUBMED 2318848
COMMENT Original source text: Homo sapiens DNA.
Drait entry and computer-readable sequence for [1] kindly submitted by A. Ichinose, 26-MAR-1990, for release after publication.
Location/Qualifiers

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Best Local Similarity 95.5%; Pred. No. 4.6e-47;
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QY      61 CACACCAAGAGTGAAGTCTGAATCTCATGTTCTGAGAAATGAGAGTCTAGGCTGTTCT 120
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QY      121 TGGAGCCACACAGAGATATGCTTGTCTAAGCTAAGCAGAGTCTGCTCACTGTG 180
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QY      181 GTCTTACCCACGCTGTGTA 201
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DB      199 GTCTTACCCACGCTGTGTA 219
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VERSION     AY192161.1 GI:27228744
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ORGANISM    Homo sapiens
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            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Homiidae; Homo
            1 (bases 1 to 55266)
            Rieder,M.J., Arnel,T.Z., Carrington,D.P., Ozuna,M., Kuldanek,S.A.,
            Rajkumar,N., Torch,E.J., Yi,Q. and Nickerson,D.A.
            Direct Submission
            Submitted (07-DEC-2002) Genome Sciences, University of Washington,
            1705 NE Pacific, Seattle, WA 98195, USA
            To cite this work please use: SeattleSNP, NHLBI HL66682 Program
            for Genomic Applications, UW-PHRC, Seattle, WA (URL:
            http://pga.gs.washington.edu).
            Location/Qualifiers
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Best Local Similarity 95.5%; Pred. No. 3.7e-47;
Matches 192; Conservative 1; Mismatches 8; Indels 0; Gaps 0;
QY 1 TTGACATCTCATCTTTCTAGGTCCTCAAGCCTTCATCTCAAGGTCATCTGGGTG 60
Db 39282 TTGACATCTCATCTTTCTAGGTCCTCAAGCCTTCATCTCAAGGTCATCTGGGTG 39341
QY 61 CACACCAAGAGTGAACCTCGAATTCATGTCAGAAATGAAAGTCTTAGGCTGTCT 120
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QY 181 GTCTTCAACCCACGCTGTGA 201
Db 39462 GTCTTCAACCCACGCTGTGA 39482
RESULT 5
HSDJ81D8 163599 bp DNA linear PRI 18-MAY-2005
LOCUS Human DNA sequence from clone RP1-81D8 on chromosome 6q25.3-26
DEFINITION Contains the PLG gene for plasminogen, the 5' end of the LPA gene
for lipoprotein (Lp(a)), a novel gene, a pseudogene similar to part
of plasminogen and a CpG island, complete sequence.
ACCESSION AL109933.25 GI:11344445
VERSION 1
KEYWORDS HTG; apolipoprotein; Cpg; kringie; lipoprotein; LPA; plasminogen;
PLG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homidae; Homo.
1 (bases 1 to 163599)
REFERENCE Tracey,A.
AUTHORS Direct Submission
TITLE Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton,
JOURNAL Cambridgeshire, CB10 1SA, UK. E-mail enquiries: veg@sanger.ac.uk
COMMENT On Nov 25, 2000 this sequence version replaced gi:11322966.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em, EMBL; Sw, SWISSPROT; Tr, TRMBL; Wp, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr6
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
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Web site: <http://www.sanger.ac.uk>
Contact: vegas@sanger.ac.uk

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC.

RPI-81D8 is from the library RPI-1 constructed by the group of Plietser de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

VECTOR: pCYPAC2

FEATURES

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VAPPTVTPVPSLEAPSEQAPTEORPGVQECYHNGOSYRGYSTVYVGT
PHSRTPIQHNRRTTENYVNAQIMNYCNRPDAVAPCYTRDPGVMEYCNL
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SRTAVTPVPSLEAPSEQAPTEORPGVQECYHNGOSYRGYSTVYVGT
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TPVPSLEAPSEQAPTEORPGVQECYHNGOSYRGYSTVYVGT
PPTVTPVPSLEAPSEQAPTEORPGVQECYHNGOSYRGYSTVYVGT
DPTVPSLEAPSEQAPTEORPGVQECYHNGOSYRGYSTVYVGT
EYVNGGLTRVYCNRPDAEIPWCYTDMPVMEYCNLTQCPVTESSVLAIVASEQ
APTQSPVQDCYHNGOSYRGYSTVYVGT
NYCNRPDAEIPWCYTDMPVMEYCNLTQCPVTESSVLAIVASEQ

gene
complement(join(16194..16339,26376..26489))
/locus_tag="Rpl-81D8.4-001"
/note="match: proteins: 018783 046507 P00747 P06867 P06868 P12345 P20918 P80009 P80010 P81286 Q15146 Q29485 Q91W05 Q9R0W3"
/pseudo

CDS
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/locus_tag="Rpl-81D8.4-001"
/note="match: proteins: 018783 046507 P00747 P06867 P06868 P12345 P20918 P80009 P80010 P81286 Q15146 Q29485 Q91W05 Q9R0W3"
/pseudo

Query Match 92.8%; Score 186.2; DB 5; Length 16359;
Best Local Similarity 95.0%; Pred. No. 1,1e-46;
Matches 191; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

Qy 1 TTGACATCCTCATCTTTCTAGTCTCTCAAGGCTTCACTACAGGTCATCTGGGTG 60
Db 89012 TTGACGTCCTCATCTTTCTAGTCTCTCAAGGCTTCACTACAGGTCATCTGGGTG 89071

Qy 61 CACACCAAGAGTGAAGCTCGAATCTCATGTTACAGAAATGAAAGTGTCTAGGCTGTTCT 120
Db 89072 CACACCAAGAGTGAATCTCGAAGCTCGAATCTCATGTTACAGAAATGAAAGTGTCTAGGCTGTTCT 89131

Qy 121 TGGAGCCCAACAGACATATTCCTGCTTAAGCTAAGACGACTCCGCTACCCGTG 180
Db 89132 TGGAGCCCAACAGACATATTCCTGCTTAAGCTAAGACGACTCCGCTACCCGTG 89191

Qy 181 GTCTTACACCCACGCTGTGA 201
Db 89192 GTCTTACACCCACGCTGTGA 89212

RESULT 6
AC010893/c 177878 bp DNA linear HTG 07-JUL-2000
LOCUS Homo sapiens chromosome 6 clone RP11-480A20, WORKING DRAFT
DEFINITION
SEQUENCE, 18 unordered pieces.
AC010893
VERSION AC010893.5 GI:8568869
HTG: HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiinae; Homo.
1 (bases 1 to 177878)
Waterston,R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 177878)
Waterston,R.H.
Direct Submission
Submitted (25-SEP-1999) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108 USA
On Jun 16, 2000 this sequence version replaced gi:6850583.

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: MUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----

Center project name: H_NH0480A20
----- Summary Statistics -----
Sequencing vector: M13; 76%
Sequencing vector: plasmid; 24%
Chemistry: Dye-terminator ET; 76% of reads
Chemistry: Dye-terminator Big Dye; 24% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 165387 bases at least Q40
Consensus quality: 170279 bases at least Q30
Consensus quality: 172646 bases at least Q20
Insert size: 183000; agarose-fp
Insert size: 176178; sum-of-contigs
Quality coverage: 4.01 in Q20 bases; sum-of-contigs
Quality coverage: 4.01 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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1 1046: contig of 1046 bp in length
1047 1146: gap of unknown length
1147 2235: contig of 1089 bp in length
2236 2335: gap of unknown length
2336 3349: contig of 1014 bp in length
3350 3449: gap of unknown length
3450 5172: contig of 1723 bp in length
5173 5272: gap of unknown length
5273 7349: contig of 2077 bp in length
7350 7449: gap of unknown length
7450 9861: contig of 2412 bp in length
9862 9961: gap of unknown length
9962 13599: contig of 3638 bp in length
13600 13699: gap of unknown length
13700 16835: contig of 3136 bp in length
16836 16935: gap of unknown length
16936 21649: contig of 4714 bp in length
21650 21749: gap of unknown length
21750 28521: contig of 6772 bp in length
28522 28621: gap of unknown length
28622 36566: contig of 7935 bp in length
36567 36566: gap of unknown length
36567 45767: contig of 9111 bp in length
45768 45867: gap of unknown length
45868 59019: contig of 13152 bp in length
59020 59119: gap of unknown length
59120 75610: contig of 16491 bp in length
75611 75710: gap of unknown length
75711 93797: contig of 18087 bp in length
93798 93897: gap of unknown length
93898 116372: contig of 22475 bp in length
116373 116472: gap of unknown length
116473 144933: contig of 28461 bp in length
144934 145033: gap of unknown length
145034 177878: contig of 32845 bp in length.
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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="6"
/clone="RP11-480A20"
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1047..1146
/estimated_length=unknown
1147..2235
/note="assembly_name:Contig41"
2236..2335
/estimated_length=unknown


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misc_feature      2336..3349
                    /note="assembly_name:Contig49"
gap               3350..3449
                    /estimated_length=unknown
misc_feature      3450..5172
                    /note="assembly_name:Contig52"
gap               5173..5272
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misc_feature      5273..7349
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gap               7350..7449
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misc_feature      7450..9861
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gap               9862..9961
                    /estimated_length=unknown
misc_feature      9962..13599
                    /note="assembly_name:Contig56"
gap               13600..13699
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misc_feature      13700..16835
                    /note="assembly_name:Contig57"
gap               16836..16935
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misc_feature      16936..21649
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gap               21650..21749
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gap               28522..28621
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gap               36557..36656
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gap               45768..45867
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gap               75611..75710
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gap               93798..93897
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vector_side:right"
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ORIGIN
Query Match      92.8%; Score 186.2; DB 12; Length 177878;
Best Local Similarity 95.0%; Pred. No. 1,1e-46;
Matches 191; Conservative 1; Mismatches 9; Indels 0; Gaps 0;
QY 1 TTGACATCCTCATCTTTTCTAGTCTCTCAAGGCTTCACTCTACAAAGTCACTCTGGGTG 60
DB 171500 TTGACGTCTCATCTTTTCTAGTCTCCCAAGGCTTCACTCTACAAAGTCACTCTGGGTG 171441

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QY 61 CACACCAAGAAGTGAACCTCGAATCTCATGTTTCAGGAAATRGAAAGTCTAGGCTGTCT 120
DB 171440 CACACCAAGAAGTGAATCTCGAACCGCATGTTTCAGGAAATAGAAAGTCTAGGCTGTCT 171381
QY 121 TGGAGCCCAACAGACAGATATTTGCTTCTTAAGCTAAGCAGAGTACTGGCTACCGTGTG 180
DB 171380 TGGAGCCCAACAGACAGAAAGATATTTGCTTCTTAAGCTAAGCAGAGTACTGGCTACCGTGTG 171321
QY 181 GTCTTACCCCAACGCTGTGA 201
DB 171320 GTCTTACCCCAACGCTGTGA 171300

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RESULT 7
AC084862/c 179154 bp DNA linear HTG 26-JUL-2005
LOCUS Papio anubis clone rp41-286p17, WORKING DRAFT SEQUENCE, 2 ordered
DEFINITION
ACCESSION AC084862
VERSION AC084862.14 GI:59797159
KEYWORDS HTG: HTGS PHASE2; HTGS DRAFT.
SOURCE Papio anubis (olive baboon)
ORGANISM Papio anubis
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Cercopithecoidea; Cercopithecinae; Papio.

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REFERENCE
AUTHORS Lau,C., Boffelli,D. and Roe,B.A.
TITLE Papio anubis BAC Clone rp41-286p17
JOURNAL Unpublished
REFERENCE
AUTHORS Lau,C., Boffelli,D. and Roe,B.A.
TITLE Direct Submission
JOURNAL Submitted (23-NOV-2000) Department Of Chemistry And Biochemistry,
The University Of Oklahoma, 620 Parrington Oval, Room 208, Norman,
OK 73019, USA

```

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REFERENCE
AUTHORS Lau,C., Boffelli,D. and Roe,B.A.
TITLE Direct Submission
JOURNAL Submitted (26-JUL-2005) Department Of Chemistry And Biochemistry,
The University Of Oklahoma, 620 Parrington Oval, Room 208, Norman,
OK 73019, USA
COMMENT
On Feb 15, 2005 this sequence version replaced gi:55701371.
----- Genome Center
Center: Department Of Chemistry And Biochemistry
The University Of Oklahoma
Center code:UOKNOR

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* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* been provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 68162: contig of 68161 bp in length
* 68162 68261: gap of unknown length
* 68262 179154: contig of 110893 bp in length.
Location/Qualifiers
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/mol_type="genomic DNA"
/db_xref="taxon:9555"
/clone="rp41-286p17"
/clone_lib="RPCI - 41 Male (Olive) Baboon BAC Library"
68162..68261
/estimated_length=unknown

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FEATURES
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Query Match      86.7%; Score 174; DB 12; Length 179154;
gap
ORIGIN

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TITLE
JOURNAL
COMMENT

Lamazares, R., Landers, T., Lehocaky, J., Levine, R., Liu, C., Liu, G.,
McDonald, P., Margulis, N., McCarthy, M., McEwan, P., McKernan, K.,
McPheters, R., Melgrim, J., Meneses, L., Mihova, T., Mlenga, V.,
Mortow, J., Murphy, T., Naylor, J., Norman, C. H., O'Connor, T.,
O'Donnell, P., O'Neill, D., Oliver, J., Oliver, J., Peterson, K.,
Pierre, N., Pisan, C., Pollara, V., Raymond, C., Rieback, M., Riley, R.,
Rogov, P., Rothman, D., Roy, A., Santos, R., Schauer, S., Severy, P.,
Sousnez, C., Spencer, B., Strange-Thomann, N., Stojanovic, N.,
Strassner, N., Subramanian, A., Talamas, J., Teffaye, S., Theodore, J.,
Tirrell, A., Travers, M., Triggilio, J., Vassiliou, H., Viel, R., Vo, A.,
Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J.,
Zimmer, A., and Zody, M.

Direct Submission

Submitted (24-ANG-2002) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

On Aug 24, 2000 this sequence version replaced gi:9799792.

All repeats were identified using RepeatMasker:

Smith, A. F. A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www.seq.wi.mit.edu

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L7625

Center clone name: 189_O_14

* NOTE: This record contains 224 individual

* sequencing reads that have not been assembled into

* contigs. Runs of N are used to separate the reads

* and the order in which they appear is completely

* arbitrary. Low-pass sequence sampling is useful for

* identifying clones that may be gene-rich and allows

* overlap relationships among clones to be deduced.

* However, it should not be assumed that this clone

* will be sequenced to completion. In the event that

* the record is updated, the accession number will

* be preserved.

* 1

* 742 841: contig of 741 bp in length

* 842 1594: contig of 753 bp in length

* 1595 1594: gap of 100 bp

* 1695 2442: contig of 748 bp in length

* 2443 2542: gap of 100 bp

* 2543 3274: contig of 732 bp in length

* 3275 3374: gap of 100 bp

* 3375 4104: contig of 730 bp in length

* 4105 4204: gap of 100 bp

* 4205 4929: contig of 725 bp in length

* 4930 5029: gap of 100 bp

* 5030 5737: contig of 708 bp in length

* 5738 5837: gap of 100 bp

* 5838 6574: contig of 737 bp in length

* 6575 6674: gap of 100 bp

* 6675 7382: contig of 708 bp in length

* 7383 7482: gap of 100 bp

* 7483 8227: contig of 745 bp in length

* 8228 8327: gap of 100 bp

* 8328 9072: contig of 745 bp in length

* 9073 9172: gap of 100 bp

* 9173 9920: contig of 748 bp in length

* 9921 10020: gap of 100 bp

* 10021 10740: contig of 720 bp in length

* 10741 10840: gap of 100 bp

* 10841 11566: contig of 726 bp in length

* 11567 11667: gap of 100 bp

* 11667 12407: contig of 741 bp in length

* 12408 12507: gap of 100 bp

* 12508 13254: contig of 747 bp in length

* 13255 13354: gap of 100 bp

* 13355 14095: contig of 741 bp in length

* 14096 14195: gap of 100 bp

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14934 15033: gap of 100 bp

15034 15755: contig of 722 bp in length

15756 15855: gap of 100 bp

15855 16584: contig of 729 bp in length

16585 16684: gap of 100 bp

16685 17433: contig of 749 bp in length

17434 17533: gap of 100 bp

17534 18286: contig of 753 bp in length

18287 18387: gap of 100 bp

18387 19127: contig of 741 bp in length

19128 19227: gap of 100 bp

19228 19937: contig of 710 bp in length

19938 20037: gap of 100 bp

20038 20750: contig of 713 bp in length

20751 20850: gap of 100 bp

20851 21574: contig of 724 bp in length

21575 21674: gap of 100 bp

21675 22395: contig of 721 bp in length

22396 22495: gap of 100 bp

22496 23225: contig of 730 bp in length

23226 23325: gap of 100 bp

23326 24072: contig of 747 bp in length

24073 24172: gap of 100 bp

24173 24902: contig of 730 bp in length

24903 25002: gap of 100 bp

25003 25746: contig of 744 bp in length

25747 25846: gap of 100 bp

25847 26585: contig of 739 bp in length

26586 26685: gap of 100 bp

26686 27375: contig of 691 bp in length

27377 27476: gap of 100 bp

27477 28217: contig of 741 bp in length

28218 28317: gap of 100 bp

28318 29039: contig of 722 bp in length

29040 29139: gap of 100 bp

29140 29876: contig of 737 bp in length

29877 29976: gap of 100 bp

29977 30717: contig of 741 bp in length

30718 30817: gap of 100 bp

30818 31554: contig of 737 bp in length

31555 31654: gap of 100 bp

31655 32380: contig of 726 bp in length

32381 32480: gap of 100 bp

32481 33227: contig of 747 bp in length

33228 33327: gap of 100 bp

33328 34072: contig of 745 bp in length

34073 34172: gap of 100 bp

34173 34896: contig of 724 bp in length

34897 34996: gap of 100 bp

34997 35739: contig of 743 bp in length

35740 35839: gap of 100 bp

35840 36559: contig of 720 bp in length

36560 36660: gap of 100 bp

36660 37393: contig of 736 bp in length

37394 37493: gap of 100 bp

37496 38105: contig of 610 bp in length

38106 38205: gap of 100 bp

38206 38948: contig of 743 bp in length

38949 39048: gap of 100 bp

39049 39799: contig of 751 bp in length

39800 39899: gap of 100 bp

39900 40646: contig of 747 bp in length

40647 40746: gap of 100 bp

40747 41467: contig of 721 bp in length

41468 42307: contig of 740 bp in length

42308 42407: gap of 100 bp

42408 43130: contig of 723 bp in length

43131 43230: gap of 100 bp

43231 43963: contig of 733 bp in length

43964 44063: gap of 100 bp

44064 44771: contig of 708 bp in length

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	Best Local Similarity	99.4%; Pred. No. 7e-37;
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Dd	45429 TTGCATCTCTCATCTTTTCATAGGTCTCAAGGCTTCATCTCACAAGTGATCTCTGGGTG	4548
Oy	61 CACCACGAAGTGAACCTCGAATCTCATGTTCAGGAATGAAGTGTCTAGGGCTGTCT	120
Dd	45489 CACACCAAGAAGTGAACCTCGAATCTCATGTTCAGGAATGAAGTGTCTAGGGCTGTCT	4554
Oy	121 TGGAGCCACACACAGACAGATAATTCGCTTGTCTAAAG	155
Dd	45549 TGGAGCCACACACAGACAGATAATTCGCTTGTCTAAAG	45583
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DEFINITION	LIPID-ASSOCIATED MOLECULES.	
VERSION	DD181590.1 GI:83947781	
KEYWORDS	JP 2005514007-A/10.	
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrate; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.	
REFERENCE	1 (bases 1 to 13869)	
AUTHORS	Bhatia,U., Tang,T.Y., Elliott,V.S., Tran,U.K., Lee,S.Y., Chang,H., Griffin,J.A., Zheng,W., Blake,J.U., Emertling,B.M., Lee,E.A., Foreythe,I.J., Bulloch,S.A., Lee,S., Reddy,R., Khare,R., Chawla,N.K., Baughn,M.R., Burrill,J.D., Ho,A. and Warren,B.A.	
TITLE	LIPID-ASSOCIATED MOLECULES	
JOURNAL	Patent: JP 2005514007-A 10-19-MAY-2005; INCYTE GENOMICS INC, Bridget A WARREN, Brooke M EMERLING, Ernestine A LEE, Hsin-Ru CHANG, Ian J FORSTYHE, Jennifer A GRIFFIN, Mariah R BAUGHN, Narinder K CHAWLA, Reena KHARE, Roopa REDDY, Sally LEE, Sean A BULLOCH, Soo Yeun LEE, Uyen K TRAN, Vicki S ELLIOTT, Tom Y TANG, Umesh BHATTIA, John D BURRILL, Julie K BLAKE, Anne HO, Wenjin ZHENG	
COMMENT	OS Homo sapiens PN JP 2005514007-A/10 PD 19-MAY-2005 PF 19-SEP-2002 JP 2003529924 PR 21-SEP-2001 US 60/324039, 02-NOV-2001 US 60/343876, PR 26-OCT-2001 US 60/346197, 14-DEC-2001 US 60/340223, PR 18-DEC-2001 US 60/342166, 22-JAN-2002 US 60/351562, PR 30-NOV-2001 US 60/334211, 03-MAY-2002 US 60/377576 PI umesh bhatia,tom y tang,vicki s elliot,uyen k tran,soo yeun lee,jennifer a griffin,wenjin zheng,julie k lee,pi hsin-ru chang,jennifer a griffin,mariah r baughn,narinder k chawla,reena khare,roopa reddy,sally lee,sean a pi brooke m emertling,erestine a lee,iان j forsythe,jean a pi bulloch,sally lee, PI roopa reddy,reena khare,narinder k chawla,mariah pi baughn,john burrill, PI d burrill, PI anne ho,bridget a warren CC This description about <220> can't be interpreted CC <220> CC <221> misc. feature CC <223> incyte ID No.: 3187560CB1 FH Key Location/Qualifiers Location/Qualifiers 1..13869 /organism="Homo sapiens" /mol_type="unassigned DNA" /db_xref="taxon:9606"	
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SOURCE		
ORIGIN		

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Dd	13150	CATGCTTAGAAGAAGTCTCTCAAGCCCTTCATCTCACAAGGTATCTGGTGACACCAA	1320	
Oy	69	GAAGTGAACCTCGAAATTCTCATGTTTCAGAGAAATGAATGTCCTAGGCCTTGAGACCC	128	
Dd	13210	GAACTGAACCTCGAATCTCATGTTTCAGAGAAATGAATGTCCTAGGCCTTGAGACCC	1326	
Oy	129	ACACAAGCAGATATTCCTGCTTAAGCTAAGCAGGTACTCGCTCACCTGTG	180	
Dd	13270	ACACAAGCAGATATTCCTGCTTAAGCTAAGCAGGTACTCGCTCACCTGTG	13321	
RESULT 11				
HSALIPA				
LOCUS	13938 bp	mRNA	linear	PRI 30-MAR-1995
DEFINITION	Human mRNA for apolipoprotein(a).			
ACCESSION	X06290	M17399	X06696	
VERSION	X06290.1	GI:28619		
KEYWORDS	apolipoprotein; apolipoprotein A; glycoprotein; lipoprotein; plasmogen; serine protease.			
SOURCE	Homo sapiens (human)			
ORGANISM	Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominoidea; Homo. 1 (bases 1 to 13938) McLean,J.W., Tomlinson,J.E., Kuang,W.J., Baton,D.L., Chen,E.Y., Fless,G.M., Scann,A.M. and Lawn,R.M. cDNA sequence of human apolipoprotein(a) is homologous to plasmogen Nature 330 (6144), 132-137 (1987)			
REFERENCE	AUTHORS McLean J.W. 2 (bases 1 to 13938) Direct Submission Submitted (23-NOV-1987) McLean J.W., Genentech Inc, 460 Point San Bruno Blvd, So. San Francisco CA 94080			
JOURNAL	TITLE Location/Qualifiers			
REFERENCE	PUBMED 3670400			
AUTHORS	McLean J.W.			
TITLE	Direct Submission			
JOURNAL	Submitted (23-NOV-1987) McLean J.W., Genentech Inc, 460 Point San Bruno Blvd, So. San Francisco CA 94080			
FEATURES	source			
CDS	1..13938 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /tissue_type="liver" /clone_id="lambda gt10, lambda ZAP" 46..13692 /note="unnamed protein product; apolipoprotein (a) (AA -19 to 4529) " /codon_start=1 /protein_id="CAA29618.1" /db_xref="GI:28620" /db_xref="GOA:P08519" /translat="MEHKYVLILLFLKSAAPQSIVDVCYHGDSYRGYSTTVT TGRTCAQMSMTPHQHNRKTENBYNLCNPDAVAAPCYTRDPGRWECNL TGCDADGTAAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVG RTCAQMSMTPHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTO CSDAEGTAAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRT COASMSMTPHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTOCS DAESTAAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRTCC AMSMTPHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTOCSDA EGTAAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRTCCAM SSMTPHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTOCSDAEG TAAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRTCCAMSS MTPHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTOCSDAEGTA VAPPTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRTCCAMSSMT PHSHSRTPREYYPNAGLMNYCRNDVAAPCYTRDPGRWECNLTOCSDAEGTAAP PTVPVPSLEAPSEQAFTPEORPGVECHNGSGYRGYSTTVGRTCCAMSSMTPH			

* consists of 7 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
* 1 10181: contig of 10181 bp in length
* 10182: gap of unknown length
* 10282: contig of 25581 bp in length
* 35863: gap of unknown length
* 35963: contig of 19466 bp in length
* 55429: gap of unknown length
* 55529: contig of 8796 bp in length
* 64325: gap of unknown length
* 77653: contig of 13228 bp in length
* 77753: gap of unknown length
* 84125: contig of 6372 bp in length
* 84225: gap of unknown length
* 150377: contig of 66153 bp in length.
Location/Qualifiers

FEATURES

source

/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
10182. 10281
/estimated_length=unknown
35863. 35963
/estimated_length=unknown
55429. 55529
/estimated_length=unknown
64325. 64424
/estimated_length=unknown
77653. 77753
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84125. 84224
/estimated_length=unknown

ORIGIN

Query Match 71.9%; Score 144.2; DB 12; Length 150377;

Best Local Similarity 85.2%; Pred. No. 1.2e-33; Indels 0; Gaps 0;

Matches 161; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

Qy 13 TCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCAAGAAG 72

Db 59197 TCTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCAAGAAG 59138

Qy 73 TGAACCTCGATCTCATGTTCAAGAAATGAAAGTCTTAGGCTGTTCTTGAGCCCAAC 132

Db 59137 AGAATCTCCAAATCAAGATGTTCAAGAAATGAAAGTCTTAGGCTGTTCTTGAGCCCAAC 59078

Qy 133 AAGCAATATTCCTGCTTAAGGCTCAAGGTCATCTGCTCAACGCTGCTTCAACCCCA 192

Db 59077 GAGCAATATTCCTGCTTAAGGCTCAAGGTCATCTGCTCAACGCTGCTTCAACCCCA 59018

Qy 193 CGCTGGTGA 201

Db 59017 AGCTGGTGA 59009

RESULT 13

AX815991

LOCUS AX815991 744 bp DNA linear PAT 09-DEC-2003

DEFINITION Sequence 64 from Patent WO03066842.

ACCESSION AX815991

VERSION AX815991.1 GI:39646630

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominoidea; Homo.

REFERENCE
AUTHORS Susilo, R., Korting, H.C., Gassen, H.G., Hils, M. and Pasternack, R.
TITLE Method for producing recombinant proteins in micro-organisms
JOURNAL Patent: WO 03066842-A 64 14-AUG-2003;
Trommsdorff GmbH & Co. KG Arzneimittel (DE)
location/Qualifiers

FEATURES

source

1. 744
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN

Query Match 66.1%; Score 132.6; DB 2; Length 744;

Best Local Similarity 85.0%; Pred. No. 6.1e-30; Indels 0; Gaps 0;

Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

Qy 8 CCTCATCTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCA 67

Db 174 CCACTGCTTGAAGAGTCCCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCA 233

Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTCTTAGGCTGTTCTTGAGACC 127

Db 234 AGAAGTGAATCTCGAATCTCATGTTCAAGAAATGAAAGTCTTAGGCTGTTCTTGAGACC 293

Qy 128 CACCAAGCAGATTTGCTTCTGCTTAAGGTCATGAGGTCATCTGCTCACTCTG 180

Db 294 CACCAAGCAGATTTGCTTCTGCTTAAGGTCATGAGGTCATCTGCTCACTCTG 346

RESULT 14

AX463624

LOCUS AX463624 750 bp DNA linear PAT 15-JUL-2002

DEFINITION Sequence 3 from Patent WO0250290.

ACCESSION AX463624

VERSION AX463624.1 GI:21886384

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominoidea; Homo.

REFERENCE
AUTHORS Negai, N., Larocque, Y. and Collen, D.J.
TITLE A yeast expression vector and a method of making a recombinant protein by expression in a yeast cell
JOURNAL Patent: WO 0250290-A 3 27-JUN-2002;
THROMB X NV (BE)

FEATURES

source

1. 750
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
<1. 750
/note="unassigned protein product"
/codon_start=1
/protein_id="CAD42144.1"
/db_xref="GI:21886385"
/translation="APSFDCGKQVEBPCKCPGRVVGCVAAHPHSMVQVSLRTPGKH
FCGGTILSPDWVLTAAHCLKSPSPSKYKVLGHNHVNLEPHVQELVSLRTPGKH
KDIALLKSSPAVITDRIYIPACLPSPHYVADRTECPITGMEGTGFFGAGLLEAOL
PVIENKCNRYEFLNGRVOSTELCAHGLAGTDSQQDSGGPLVCFEKKYILGVTIS
WGLGCAKPNKPGYVYVRSRFTWIEGMNRN"

CDS

Qy 8 CCTCATCTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCA 67

Db 180 CCACTGCTTGAAGAGTCCCAAGGCTTCATCTCAAGGTCATCTGAGTGCACACCA 239

Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTCTTAGGCTGTTCTTGAGACC 127

ORIGIN

Query Match

Best Local Similarity 66.1%; Score 132.6; DB 2; Length 750;

Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

Db 240 AGAAGTGAATCTCGAACCGCAGATTCAGAAATAGAGTCTAGGCTGTCTTGAGACC 299
Oy 128 CACACAAGCAGATATTGCTTGTCTTAAGTAAGCAGAGTACTGCTCACCTGTG 180
Db 300 CACACGAAAAGATATTGCTTGTCTTAAGTAAGCAGAGTACTGCTCACCTGTG 352

RESULT 15

AX815990

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

Homnidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

FEATURES

source

1. 750

/organism="Homo sapiens"

/mol_type="unassigned DNA"

/db_xref="taxon:9606"

ORIGIN

Query Match

Best Local Similarity

Matches

147; Conservative

1; Mismatches

25; Indels

0; Gaps

0;

Oy

Db

Oy

Db

Oy

Db

Oy

Db

Oy

Db

Oy

Db

Oy

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Db

Oy

Db

Search completed: May 26, 2006, 15:57:00
Job time : 2038 secs

Query Match	66.9%;	Score 134.2;	DB 14;	Length 2141;
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IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1004 Std Error: 0.00
Seq primer: M13R1
High quality sequence stop: 293.
Location/Qualifiers

FEATURES
source

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1. .459
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/mol_type="mRNA"
/db_xref="GDB:3767398"
/db_xref="taxon:9606"
/clone="IMAGE:198348"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/clone_idb="Soares fetal liver spleen INFLS"
/notes="Organ: Liver and Spleen; Vector: pT73D (Pharmacia)
with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo(dT) primer
[5' AACGGAAGATTAATTAAGATCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
```

ORIGIN

Query Match 66.1%; Score 132.6; DB 10; Length 459;
Best Local Similarity 85.0%; Pred. No. 2.1e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

Qy 8 CCTATCTTTTCTAGTCTCTCAAGGCTTCTATCTTCAAGATCCTCGGGTGCACACCA 67
Db 54 CCATGCTTGGAGAGTCCCAAGGCTTCTATCTTCAAGATCCTCGGGTGCACACCA 113
Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAGATGCTAGGCTGTTCTTGAGACC 127
Db 114 AGAAGTGAATCTCGAACCGCATGTTCAAGAAATGAGATGCTAGGCTGTTCTTGAGACC 173
Qy 128 CACACAGACAGATATTTGCTTCTTAAGCTAAGCAAGCACTGCTCAGCTGTG 180
Db 174 CACACGAAAGATATTTGCTTCTTAAGCTAAGCAAGCTGCTCAGCTGTG 226

RESULT 6
A1948806/c 466 bp mRNA linear EST 08-MAR-2000
LOCUS WQ26612.x1 NCI CGAP Kid11 Homo sapiens cDNA clone IMAGE:2472430.3
DEFINITION similar to gb:X05199 PLASMINOEN PRECURSOR (HUMAN); mRNA sequence.
ACCESSION A1948806
VERSION A1948806.1 GI:5741116
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.
1 (bases 1 to 466)
NCI-CCGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA library Preparation: M. Bento Soares, Ph.D.
CDNA library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CCGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 535 Std Error: 0.00
Seq primer: -40UP from Gibco

High quality sequence stop: 430.
Location/Qualifiers

FEATURES
source

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/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2472430"
/lab_host="DH10B"
/clone_idb="NCI CGAP Kid11"
/notes="Organ: Kidney; Vector: pT73D-PacI; Site 1: Not I;
Site 2: Eco RI; Plasmid DNA from the normalized library
NCI CGAP Kid3 was prepared, and ss circles were made in
vitro. Following Hsp purification, this DNA was used as
tracer in a subtractive hybridization reaction. The driver
was PCR-amplified cDNAs from a pool of 5,000 clones made
from the same library (cloneids 132376-132391).
145007-1456775, and 1500552-1502855). Subtraction by
Bento Soares and M. Fatima Bonaldo."
```

ORIGIN

Query Match 66.1%; Score 132.6; DB 1; Length 466;
Best Local Similarity 85.0%; Pred. No. 2.1e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

Qy 8 CCTATCTTTTCTAGTCTCTCAAGGCTTCTATCTTCAAGATCCTCGGGTGCACACCA 67
Db 246 CCATGCTTGGAGAGTCCCAAGGCTTCTATCTTCAAGATCCTCGGGTGCACACCA 187
Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAGATGCTAGGCTGTTCTTGAGACC 127
Db 186 AGAAGTGAATCTCGAACCGCATGTTCAAGAAATGAGATGCTAGGCTGTTCTTGAGACC 127
Qy 128 CACACAGACAGATATTTGCTTCTTAAGCTAAGCAAGCACTGCTCAGCTGTG 180
Db 126 CACACGAAAGATATTTGCTTCTTAAGCTAAGCAAGCTGCTCAGCTGTG 74

RESULT 7

N91337 475 bp mRNA linear EST 03-APR-1996
LOCUS za14h05.x1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
DEFINITION IMAGE:292569.5' similar to gb:X05199 PLASMINOEN PRECURSOR
(HUMAN); mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

N91337
N91337.1 GI:1444664
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.
1 (bases 1 to 475)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,
Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marx, M.,
Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
Trevaskis, E., Waterston, R., Williamson, A., Wohlmann, P. and
Wilson, R.

REFERENCE
AUTHORS
JOURNAL
COMMENT

The Mashu-Merck EST Project
Unpublished (1995)
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: mob.BEBA+R
High quality sequence stop: 296.
Location/Qualifiers
1. .475
/organism="Homo sapiens"
/mol_type="mRNA"

RESULT 12
AV661991
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AV661991
AV661991 GLC Homo sapiens cDNA clone GLC2G06 3', mRNA sequence.
AV661991
AV661991.1 GI:9883005
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
PUBMED
COMMENT

1 (bases 1 to 594)
Xu, X., Huang, J., Xu, Z., Qian, B., Zhu, Z., Yan, Q., Cai, T., Zhang, X.,
Xiao, H., Qu, J., Liu, F., Huang, Q., Cheng, Z., Li, N., Du, J., Hu, W.,
Shen, K., Lu, G., Fu, G., Zhong, M., Xu, S., Gu, W., Huang, W., Zhao, X.,
Hu, G., Gu, J., Chen, Z., and Han, Z.
Insight into hepatocellular carcinogenesis at transcriptome level
by comparing gene expression profiles of hepatocellular carcinoma
with those of corresponding noncancerous liver
Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15089-15094 (2001)
11752456
Contact: Zeguang Han
Chinese National Human Genome Center at Shanghai
351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
201203, P. R. China
Tel: 86-21-50801919 (ex. 45)
Fax: 86-21-50801922
Email: hanzg@chgc.sh.cn
This clone is available at CHGC in Shanghai.

FEATURES
Source
Location/Qualifiers

1..594
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/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="GLC2G06"
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/dev_stage="Adult"
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/note="Vector: pBluescript sk(-); Site_1: EcoRI; Site_2:
XhoI"

ORIGIN

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Best Local Similarity 85.0%; Pred. No. 2.3e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTTCTAGTCTCTCAAGGCTTCATCTCAAGGTCATCTGGTGACACCA 67
DB 123 CCACGTGCTTGAGAAATCCCAAGGCTTCATCTCAAGGTCATCTGGTGACACCA 182
QY 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGACC 127
DB 183 AGAAGTGAATCTCGAACCAGCATTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGACC 242
QY 128 CACACAGAGATATTTGCTTGTCTAAAGCTAAGCAGTACTGCTCACTGTG 180
DB 243 CACACAGAAAGATATTTGCTTGTCTAAAGCTAAGCAGTACTGCTCACTGTG 295

RESULT 13
AV662084
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AV662084
AV662084 GLC Homo sapiens cDNA clone GLCHAG06 3', mRNA sequence.
AV662084
AV662084.1 GI:9883098
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.
1 (bases 1 to 611)
Xu, X., Huang, J., Xu, Z., Qian, B., Zhu, Z., Yan, Q., Cai, T., Zhang, X.,
Xiao, H., Qu, J., Liu, F., Huang, Q., Cheng, Z., Li, N., Du, J., Hu, W.,
Shen, K., Lu, G., Fu, G., Zhong, M., Xu, S., Gu, W., Huang, W., Zhao, X.,
Hu, G., Gu, J., Chen, Z., and Han, Z.
Insight into hepatocellular carcinogenesis at transcriptome level
by comparing gene expression profiles of hepatocellular carcinoma
with those of corresponding noncancerous liver
Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15089-15094 (2001)
11752456
Contact: Zeguang Han
Chinese National Human Genome Center at Shanghai
351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
201203, P. R. China
Tel: 86-21-50801919 (ex. 45)
Fax: 86-21-50801922
Email: hanzg@chgc.sh.cn
This clone is available at CHGC in Shanghai.

FEATURES
Source
Location/Qualifiers

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/organism="Homo sapiens"
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/note="Vector: pBluescript sk(-); Site_1: EcoRI; Site_2:
XhoI"

ORIGIN

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Best Local Similarity 85.0%; Pred. No. 2.3e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTTCTAGTCTCTCAAGGCTTCATCTCAAGGTCATCTGGTGACACCA 67
DB 122 CCACGTGCTTGAGAAATCCCAAGGCTTCATCTCAAGGTCATCTGGTGACACCA 181
QY 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGACC 127
DB 182 AGAAGTGAATCTCGAACCAGCATTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGACC 241
QY 128 CACACAGAGATATTTGCTTGTCTAAAGCTAAGCAGTACTGCTCACTGTG 180
DB 242 CACACAGAAAGATATTTGCTTGTCTAAAGCTAAGCAGTACTGCTCACTGTG 294

RESULT 14
B1759134
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

B1759134
B1759134 NIH_MGC_116 Homo sapiens cDNA clone IMAGE:5183003 5',
mRNA sequence.
B1759134
B1759134.1 GI:15750712
EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

NIH-MGC http://mhc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cga@bbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov

Place: LLNL1456 row: 1 column: 12
High quality sequence stop: 766.
Location/Qualifiers

FEATURES
source

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/clone_1ib="NIH_MGC_116"
/note="Organ: pooled colon, kidney, stomach; Vector:
pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA
source anonymous pool of 3 colons, age 26 yo male, 49 yo
female, 71 yo male colon; 46 yo male kidney, and pool of 2
stomachs, 62 yo male and 70 yo female. Library is
oligo-dT primed and directionally cloned (EcoRV site is
destroyed upon cloning). Average insert size 1.4 kb.
Insert size range 1-3 kb. Library is normalized and
enriched for full-length clones and was constructed by C.
Gruber (Invitrogen). Research Genetics tracking code
023. Note: this is a NIH_MGC Library."
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ORIGIN

```
Query Match      66.1%; Score 132.6; DB 2; Length 766;
Best Local Similarity 85.0%; Pred. No. 2.4e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTTCTAGAGCTTCAAGGCTTCATCTCAAGGTCAATCCTGGTGACACCA 67
DB 258 CCACGCTTGGAGAAAGTCCCAAGGCTTCATCTCAAGGTCAATCCTGGTGACACCA 317
QY 68 AGAAGTGAACCTCGAATCTCATGTTCAGAAATGGAAGTGTAGGCTGTTTGGAGCC 127
DB 318 AGAAGTGAATCTCGAAGCCGATGTTCAAGAAATAGAGTGTAGGCTGTTTGGAGCC 377
QY 128 CACACAGCAGATATTTGCTTCTTAAGCTAAGCAGTACTGCTCACCTGTG 180
DB 378 CACAGCAAAAGATATTGCTTCTGTAAAGCTAAGCAGTACTGCTCACCTGTG 430
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RESULT 15

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LOCUS      BG402207      859 bp      mRNA      linear      EST 12-MAR-2001
DEFINITION 602465764F1 NIH_MGC_75 Homo sapiens cDNA IMAGE:4594023 5',
            mRNA sequence.
ACCESSION  BG402207
VERSION    BG402207.1 GI:13295655
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Homnidae; Homo.
REFERENCE  1 (bases 1 to 859)
            NIH-MGC http://mgs.nci.nih.gov/.
            National Institutes of Health, Mammalian Gene Collection (MGC)
            Unpublished (1999)
            Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: CLONTECH Laboratories, Inc.
            cDNA Library Preparation: CLONTECH Laboratories, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: L10M1334 row: 0 column: 16
            High quality sequence stop: 629.
            Location/Qualifiers
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FEATURES

source

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1. 859
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4594023"
/lab_host="DH10B (T1 phage-resistant)"
/clone_1ib="NIH_MGC_75"
/note="Organ: kidney; Vector: pDNR-LIB (Clontech); Site 1:
SfiI (ggcgccgcgcgc); Site 2: SfiI (ggccatcatggcc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-ATCTAGAGGCGGAGGCGGCGGACATG-3' and 3' adaptor sequence:
5'-ATTCTAGAGGCGGAGGCGGCGGACATG-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size 1.65
kb (range 0.5-4.0 kb). 15/15 colonies contained inserts
by PCR. This library was enriched for full-length clones
and was constructed by Clontech Laboratories (Palo Alto,
CA). Note: this is a NIH_MGC Library."
```

ORIGIN

```
Query Match      66.1%; Score 132.6; DB 2; Length 859;
Best Local Similarity 85.0%; Pred. No. 2.5e-31;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTTCTAGAGCTTCTCAAGGCTTCATCTCAAGGTCAATCCTGGTGACACCA 67
DB 313 CCACGCTTGGAGAAAGTCCCAAGGCTTCATCTCAAGGTCAATCCTGGTGACACCA 372
QY 68 AGAAGTGAACCTCGAATCTCATGTTCAGAAATGGAAGTGTAGGCTGTTTGGAGCC 127
DB 373 AGAAGTGAATCTCGAAGCCGATGTTCAAGAAATAGAGTGTAGGCTGTTTGGAGCC 432
QY 128 CACACAGCAGATATTTGCTTCTTAAGCTAAGCAGTACTGCTCACCTGTG 180
DB 433 CACAGCAAAAGATATTGCTTCTGTAAAGCTAAGCAGTACTGCTCACCTGTG 485
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Search completed: May 26, 2006, 16:10:41
Job time: 4490 secs

GenCore version 5.1.8
Copyright (c) 1993 - 2006 Bioacceleration Ltd.

OM nucleic - nucleic search, using bw model

Run on: May 26, 2006, 12:35:58 ; Search time 354 Seconds
(without alignments)
3958.819 Million cell updates/sec

Title: US-10-796-280-19350
Perfect score: 200.6
Sequence: 1 tctgacctccctccatcttctc.....tcttcacccacgcgtgtga 201

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 5244920 seqs, 3486124231 residues

Total number of hits satisfying chosen parameters: 10489840

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing filter 45 summaries

Database :
1: Genesegq1980s:*
2: Genesegq1990s:*
3: Genesegq2000s:*
4: Genesegq2001as:*
5: Genesegq2001bs:*
6: Genesegq2002as:*
7: Genesegq2002bs:*
8: Genesegq2003as:*
9: Genesegq2003bs:*
10: Genesegq2003cs:*
11: Genesegq2003ds:*
12: Genesegq2004as:*
13: Genesegq2004bs:*
14: Genesegq2005s:*
15: Genesegq2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	144.4	72.0	201	13 ADO45134	Adq45134 Myocardia
2	144.4	72.0	201	13 ADO45152	Adq45152 Myocardia
3	144.4	72.0	820	5 AAS80437	Aas80437 DNA encod
4	144.4	72.0	820	5 AAS66111	Aas66111 DNA encod
5	144.4	72.0	3756	13 ADO38544	Adq38544 Human SNP
6	144.4	72.0	10422	8 ACA64741	Ac64741 Apolipop
7	144.4	72.0	10422	10 ADF43405	Adf43405 Apolipop
8	144.4	72.0	10422	14 AEA03076	Aea03076 iNOS nucl
9	144.4	72.0	13869	8 ACC47280	Acc47280 Human lip
10	144.4	72.0	13938	8 ACC47280	Acc47280 Human apo
11	144.4	72.0	13938	12 ADO3412	Ado3412 Human apo
12	144.4	72.0	13938	12 ADO75139	Ado75139 Human apo
13	144.4	72.0	13938	13 ADO38545	Ado38545 Human SNP
14	144.4	72.0	2178	5 AAS80436	Aas80436 DNA encod
15	132.6	66.1	744	10 ACH00783	Ach00783 Human mic
16	132.6	66.1	750	6 ABR89460	Ab89460 Human mic
17	132.6	66.1	750	6 ACH00782	Ach00782 Human mic
18	132.6	66.1	750	13 ADS20373	Ads20373 Human mic

19	132.6	66.1	783	10 ACH00781	Ach00781 Human mic
20	132.6	66.1	786	10 ACH00780	Ach00780 Human mic
21	132.6	66.1	999	10 ACH00777	Ach00777 Human-S C
22	132.6	66.1	1005	10 ACH00775	Ach00775 Human-S C
23	132.6	66.1	1011	10 ACH00778	Ach00778 Human-S C
24	132.6	66.1	1017	10 ACH00776	Ach00776 Human-S C
25	132.6	66.1	1038	10 ACH00773	Ach00773 Human-S C
26	132.6	66.1	1041	10 ACH00771	Ach00771 Human-S C
27	132.6	66.1	1047	6 ABR89461	Ab89461 Human min
28	132.6	66.1	1047	10 ACH00779	Ach00779 Human min
29	132.6	66.1	1047	13 ADS20377	Ads20377 Human min
30	132.6	66.1	1050	10 ACH00774	Ach00774 Human-S C
31	132.6	66.1	1053	10 ACH00772	Ach00772 Human-S C
32	132.6	66.1	1302	10 ACH00769	Ach00769 Human-S C
33	132.6	66.1	1314	10 ACH00770	Ach00770 Human-S C
34	132.6	66.1	1724	2 AAQ40318	Aaq40318 Sequence
35	132.6	66.1	1907	8 ACC51086	Acc51086 Human Pla
36	132.6	66.1	2145	10 ACH00785	Ach00785 Human Pla
37	132.6	66.1	2236	2 AAQ40258	Aaq40258 Plasmid p
38	132.6	66.1	2236	3 AAB89829	Aab89829 Plasmid p
39	132.6	66.1	2296	4 AAS12747	Aas12747 Plasmid p
40	132.6	66.1	2376	10 ACH00784	Ach00784 Human Glu
41	132.6	66.1	2376	13 ADS20379	Ads20379 Human pla
42	132.6	66.1	2400	10 ACH00752	Ach00752 Human-S C
43	132.6	66.1	2412	10 ACH00753	Ach00753 Human-S C
44	132.6	66.1	2430	10 ABX15877	Abx15877 DNA encod
45	132.6	66.1	2432	14 AED76372	Aed76372 Human pla

ALIGNMENTS

RESULT 1
ADQ45134 standard; DNA; 201 BP.
ID ADQ45134
AC ADQ45134;
DT 18-NOV-2004 (first entry)
XX
DE Myocardial infarction-associated SNP flanking transcript, SEQ ID 6797.
XX
KW Myocardial infarction; detection; single nucleotide polymorphism; SNP;
XX
KW cardiant; gene therapy; human; ds.
XX
OS Homo sapiens.
XX
FN WO2004058052-A2.
XX
PD 15-JUL-2004.
XX
PP 22-DEC-2003; 2003WO-US040978.
XX
PR 20-DEC-2002; 2002US-0434778P.
PR 10-MAR-2003; 2003US-0453135P.
PR 30-APR-2003; 2003US-0466412P.
PR 23-SEP-2003; 2003US-0504955P.
XX
PA (APPL-) ABPLERA CORP.
XX
PI Cargill M, Devlin JJ, Iakubova O;
XX
DR WPI; 2004-533949/51.
XX
PT Identifying an individual who has an altered risk for developing
PT myocardial infarction by detecting a single nucleotide polymorphism in
PT the individual's nucleic acids.
XX
PS Claim 7; SEQ ID NO 6797; 145bp; English.
XX
CC The invention relates to a novel method for identifying an individual who
CC has an altered risk for developing myocardial infarction. The method
CC comprises detecting a single nucleotide polymorphism (SNP) in any one of

CC the nucleotide sequences given in the specification in the individual's
CC nucleic acid, where the presence of the SNP is correlated with an
CC altered risk for myocardial infarction in the individual. The invention
CC further comprises: an isolated nucleic acid molecule comprising at least
CC 8 contiguous nucleotides where one of the nucleotides is an SNP given in
CC the specification or its complement and encoding any one of the amino
CC acid sequences given in the specification; an isolated polypeptide
CC comprising an amino acid sequence given in the specification; an antibody
CC that specifically binds to the polypeptide or its antigen-binding
CC fragment; an amplified polynucleotide containing an SNP given in the
CC specification and which is between about 16 and 1000 nucleotides in
CC length; a kit for detecting an SNP in a nucleic acid, comprising the
CC polynucleotide, a buffer and an enzyme; a method of detecting an SNP in a
CC nucleic acid molecule; a method of detecting a variant polypeptide; and a
CC method for identifying an agent useful in treating or preventing
CC myocardial infarction. The novel detection method has cardiac activity.
CC The nucleic acids of the invention may be used in gene therapy. The
CC method is useful in identifying an individual who has an increased or
CC decreased risk for developing myocardial infarction and for preparing a
CC composition for treating or preventing myocardial infarction. This
CC polynucleotide sequence represents a transcript-based context sequence
CC which flanks an SNP found in a human myocardial infarction-associated
CC gene of the invention. Note: This sequence was not shown in the
CC specification. The sequence has come from an electronic sequence listing
CC downloaded from the WIPO website.

Sequence 201 BP; 53 A; 54 C; 45 G; 48 T; 0 U; 1 Other;

Query Match	72.0%	Score 144.4	DB 13	Length 201
Best Local Similarity	90.1%	Pred. No. 5.1e-40		
Matches 155; Conservative	0	Mismatches 17	Indels 0	Gaps 0

QY 9 CTATCTTTCTAGGCTCTCAAGGCTTATCTTCAACAAAGTATCTTGGGACACAA 68
Db 9 CACTGCTTGAAGAAGTCTTAAAGCTTATCTTCAAAAGTATCTTGGGACACAA 68
QY 69 GAACTGAACCTCGAATCTCATTTCAAGAAATGAAAGCTTAAAGCTTCTTGAAGCC 128
Db 69 GAACTGAACCTCGAATCTCATTTCAAGAAATGAAAGCTTAAAGCTTCTTGAAGCC 128
QY 129 ACACAAGCAGATATTGCTTGCTTAAAGCTTAAAGGTAAGTCTGCTCACTGTG 180
Db 129 ACACAAGCAGATATTGCTTGCTTAAAGCTTAAAGGCTGCGCTGATACAGT 180

RESULT 2
ADQ45152
ID ADQ45152 standard; DNA; 201 BP

AC	ADQ45152;
XX	
DT	18-NOV-2004 (first entry)

DE	Myocardial infarction-associated SNP flanking transcript, SEQ ID 6015
XX	
KW	Myocardial infarction; detection; single nucleotide polymorphism; SNP
KW	cardiac; gene therapy; human; ds.

OS Homo sapiens.

PN WO2004058052-A2.

PD 15-JUL-2004

Pf 22-DEC-2003; 2003WO-US040978.

PR	20-DEC-2002;	2002US-0434778P
PR	10-MAR-2003;	2003US-0453135P
PR	30-APR-2003;	2003US-0466412P
PR	23-SEP-2003;	2003US-0504955P

PA (APPL-) APPLERA CORP

PI Cargill M, Devlin JJ, Iakubova O,
 XX
 DR WPI; 2004-533949/51.
 XX
 XX
 PT Identifying an individual who has an altered risk for developing
 XX myocardial infarction by detecting a single nucleotide polymorphism in
 PT the individual's nucleic acids.

PS Claim 7; SEQ ID NO 6815; 145pp; English.

CC The invention relates to a novel method for identifying an individual who
CC has an altered risk for developing myocardial infarction. The method
CC comprises detecting a single nucleotide polymorphism (SNP) in any one of
CC the nucleotide sequences given in the specification in the individual's
CC nucleic acids, where the presence of the SNP is correlated with an
CC altered risk for myocardial infarction in the individual. The invention
CC further comprises: an isolated nucleic acid molecule comprising at least
CC 8 contiguous nucleotides where one of the nucleotides is an SNP given in
CC the specification or its complement and encoding any one of the amino
CC acid sequences given in the specification; an isolated polypeptide
CC comprising an amino acid sequence given in the specification; an antibody
CC that specifically binds to the polypeptide or its antigen-binding
CC fragment; an amplified polynucleotide containing an SNP given in the
CC specification and which is between about 16 and 1000 nucleotides in
CC length; a kit for detecting an SNP in a nucleic acid, comprising the
CC polynucleotide, a buffer and an enzyme; a method of detecting an SNP in a
CC nucleic acid molecule; a method of detecting a variant polypeptide; and a
CC method for identifying an agent useful in treating or preventing
CC myocardial infarction. The novel detection method has cardiant activity.
CC The nucleic acids of the invention may be used in gene therapy. The
CC method is useful in identifying an individual who has an increased or
CC decreased risk for developing myocardial infarction and for preparing a
CC composition for treating or preventing myocardial infarction. This
CC polynucleotide sequence represents a transcript-based context sequence
CC which flanks an SNP found in a human myocardial infarction-associated
CC gene of the invention. Note: This sequence was not shown in the
CC specification. The sequence has come from an electronic sequence listing
CC downloaded from the WIPO website.

Sequence 201 BP; 53 A; 54 C; 45 G; 48 T; 0 U; 1 Other;

Query Match	72.0%	Score 144.4	DB 13	Length 201
Best Local Similarity	90.1%	Pred. No. 5.1e-40		
Matches 155; Conservative	0	Mismatches 17	Indels 0	Gaps 0

OY	9	CTCATCTTTTCTAGSTCTCTCAAGGCGCTTCATCTTACAAAGTATCCTGGGTGACACACCA	68
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OY	69	GAAGTGAACCTTCGAATCTCATGTTTACGAAATRGAAGTCTTCAAGGCTGTTCTTGGAGCC	128
Db	69	GAAGTGAACCTTCGAATCTCATGTTTACGAAATRGAAGTCTTCAAGGCTGTTCTTGGAGCC	128
OY	129	ACACAGAGATATTGCGCTGTCTAAGCTAAGCTAACAGATCTCGCTCACTGTG	180
Db	129	ACACAGAGATATTGCGCTGTCTAAGCTAAGCTAACAGAGCTCGCGTATCACTG	180

RESULT 3
AAS80437
ID AAS80437 standard; cDNA; 820 BP.

AC AAS80437;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #16241.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; ss

Homo sapiens

FN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US008631.
 XX
 PR 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 DR P-PSDB; ABG16250.
 XX
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 PS Claim 1; SEQ ID NO 16241; 103bp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
 CC and in recombinant production of (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
 CC coding sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 CC
 XX Sequence 820 BP; 209 A; 203 C; 209 G; 199 T; 0 U; 0 Other.
 SQ
 Query Match 72.0%; Score 144.4; DB 5; Length 820;
 Best Local Similarity 89.5%; Pred. No. 9.7e-40;
 Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
 QY 9 CTCATCTTTTCTAGTCTCTCAAGGCTTCATCTACAGGTATCTGGGTGACACCA 68
 DB 116 CACTCTCTGAAGAAATCTCAAGGCTTCATCTACAGGTATCTGGGTGACACCA 175
 QY 69 GAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCCC 128
 DB 176 GAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCCC 235
 QY 129 ACACAAGAGATATGCTTGTCTAAAGCTAAGCAGGTAAGTCTGCTCACTGTG 180
 DB 236 ACACAAGAGATATGCTTGTCTAAAGCTAAGCAGGCTGCTGCTATCACTG 287
 RESULT 4
 AAS6111
 ID AAS6111 standard; cDNA; 820 BP.
 AC AAS6111;
 AC AAS6111;
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #1915.

XX
 XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US008631.
 XX
 PR 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 DR P-PSDB; ABG01924.
 XX
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 PS Claim 1; SEQ ID NO 1915; 103bp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
 CC and in recombinant production of (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
 CC coding sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 CC
 XX Sequence 820 BP; 209 A; 203 C; 209 G; 199 T; 0 U; 0 Other.
 SQ
 Query Match 72.0%; Score 144.4; DB 5; Length 820;
 Best Local Similarity 89.5%; Pred. No. 9.7e-40;
 Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
 QY 9 CTCATCTTTTCTAGTCTCTCAAGGCTTCATCTACAGGTATCTGGGTGACACCA 68
 DB 116 CACTCTCTGAAGAAATCTCAAGGCTTCATCTACAGGTATCTGGGTGACACCA 175
 QY 69 GAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCCC 128
 DB 176 GAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCCC 235
 QY 129 ACACAAGAGATATGCTTGTCTAAAGCTAAGCAGGTAAGTCTGCTCACTGTG 180
 DB 236 ACACAAGAGATATGCTTGTCTAAAGCTAAGCAGGCTGCTGCTATCACTG 287
 RESULT 5
 AD038544
 ID AD038544 standard; DNA; 3756 BP.

XX ADO38544;
AC 18-NOV-2004 (first entry)
XX
XX
DE Human SNP containing myocardial infarction-associated gene, SEQ ID 207.
XX
XX Myocardial infarction; detection; single nucleotide polymorphism; SNP;
KM cardiant; gene therapy; human; gene; ds.
XX
OS Homo sapiens.
XX MO2004058052-A2.
XX
XX 15-JUL-2004.
XX
XX 22-DEC-2003; 2003WO-US040978.
XX
XX 20-DEC-2002; 2002US-0434778P.
PR 10-MAR-2003; 2003US-0453135P.
PR 30-APR-2003; 2003US-0466412P.
PR 23-SEP-2003; 2003US-0504955P.
XX
XX (APPL-) APPLERA CORP.
XX
XX Cargill M, Devlin JJ, Iakubova O;
PI WPI; 2004-533949/51.
XX
DR P-PSDB; ADO39372.
XX
XX Identifying an individual who has an altered risk for developing
PT myocardial infarction by detecting a single nucleotide polymorphism in
PT the individual's nucleic acids.
XX
XX Claim 7, SEQ ID NO 207, 145bp; English.
XX
XX The invention relates to a novel method for identifying an individual who
CC has an altered risk for developing myocardial infarction. The method
CC comprises detecting a single nucleotide polymorphism (SNP) in any one of
CC the nucleotide sequences given in the specification in the individual's
CC nucleic acids, where the presence of the SNP is correlated with an
CC altered risk for myocardial infarction in the individual. The invention
CC further comprises: an isolated nucleic acid molecule comprising at least
CC 8 contiguous nucleotides where one of the nucleotides is an SNP given in
CC the specification or its complement and encoding any one of the amino
CC acid sequences given in the specification; an isolated polypeptide
CC comprising an amino acid sequence given in the specification; an antibody
CC that specifically binds to the polypeptide or its antigen-binding
CC fragment; an amplified polynucleotide containing an SNP given in the
CC specification and which is between about 16 and 1000 nucleotides in
CC length; a kit for detecting an SNP in a nucleic acid, comprising the
CC polynucleotide, a buffer and an enzyme; a method of detecting an SNP in a
CC nucleic acid molecule; a method of detecting a variant polypeptide; and a
CC method for identifying an agent useful in treating or preventing
CC myocardial infarction. The novel detection method has cardiant activity.
CC The nucleic acids of the invention may be used in gene therapy. The
CC method is useful in identifying an individual who has an increased or
CC decreased risk for developing myocardial infarction and for preparing a
CC composition for treating or preventing myocardial infarction. This
CC polynucleotide sequence represents a human myocardial infarction-
CC associated gene containing one or more SNPs of the invention. Note: This
CC sequence was not shown in the specification. The sequence has come from
CC an electronic sequence listing downloaded from the WIPO website.
XX
SO Sequence 3756 BP, 1017 A; 965 C; 914 G; 839 T; 0 U; 21 Other;

Query Match 72.0%; Score 144.4; DB 13; Length 3756;
Best Local Similarity 90.1%; Pred. No. 1.9e-39;
Matches 155; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

9 CTGATCTTTTAGTCTCTCAAGGCTTCATCTCTCAAGGTCATCTCTGAGTGCACACCAA 68
DB CACTGCTTGAAGAGTCTCTCAAGGCTTCATCTCTCAAGGTCATCTCTGAGTGCACACCAA 3027

QY 69 GAAGTGAACCTCGAATCTCATGTTACAGAAATRGAACTGTCTAGCTGTTCTTGAGCC 128
DB 3028 GAAGTGAACCTCGAATCTCATGTTACAGAAATRGAACTGTCTAGCTGTTCTTGAGCC 3087
QY 129 ACACACACAGATATTTGCTTGTCTTAAAGCTAAGCAGGCTACCTGCTGAGCTG 180
DB 3088 ACACACACAGATATTTGCTTGTCTTAAAGCTAAGCAGGCTGCTGCTATCATCTG 3139

RESULT 6
ACA64741
ID ACA64741 standard; DNA; 10422 BP.
XX
XX ACA64741;
XX
XX 18-JUN-2003 (first entry)
XX
DE Apolipoprotein gene #1.
XX
XX Superantigen; ds; gene; SAg; staphylococcal enterotoxin; tumour; cancer;
KM apoptosis; gene therapy; mammalian cell receptor; cytosolic;
KM tumour associated lipid; energy; T cell; antigen presenting cell; APC;
KM tumouricidal immunocyte; antitumour.
XX
XX Unidentified.
XX
XX US2002177551-A1.
XX
XX 28-NOV-2002.
XX
XX 30-MAY-2001; 2001US-00870759.
XX
PR 31-MAY-2000; 2000US-0208128P.
XX
XX (TERM/) TERMAN D S.
XX
XX Terman DS;
XX
XX WPI; 2003-361759/34.
XX
DR P-PSDB; ABU79139.
XX
XX A mammalian cell receptor, useful in the treatment of cancer by binding
PT to tumour associated lipids where the binding induces energy or apoptosis
PT in T cells and antigen presenting cells.
XX
XX Example 2; Page; 167pp; English.
XX
XX The invention relates to a mammalian cell receptor, useful in the
CC treatment of cancer, which binds to tumour associated lipid and induces
CC energy or apoptosis in the T cells and antigen presenting cells (APCs).
CC Also included are a mammalian cell useful in the treatment of cancer
CC where the receptor which binds tumour associated lipids and induces
CC cellular inactivation or death is deleted or functionally deactivated,
CC producing (M1) a tumouricidal immunocyte population in vivo in a mammal
CC (by allowing tumour suppressive fatty acids, ceramides, glycolipids,
CC receptors for immunosuppressive fatty acids, gangliosides, sphingolipids,
CC sphingolipids, glycosphingolipids, phosphosphingolipids, gangliosides,
CC sialylated glycans, lipopeptides and proteoglycolipids are inactivated or
CC deleted) a construct useful in the treatment of cancer comprising a
CC superantigen (SAg) nucleotide inserted into a virus, a mammalian T cell
CC useful in the treatment of cancer (where an adaptor protein in which
CC inhibits T cell activation by tumour associated antigens is deleted or
CC functionally deactivated), a composition useful in the treatment of
CC cancer (comprising a lipid raft conjugated to a superantigen), producing
CC (M2) a tumouricidal immunocyte population ex vivo in a mammal (by
CC allowing tumour associated lipids to contact immunocytes, in which
CC receptors for the lipids are inactivated or deleted to produce a
CC tumouricidal immunocyte population, and administering the tumouricidally
CC activated immunocytes to the host), producing (M3) a tumouricidal APC
CC population ex vivo in a mammal (by allowing a tumour associated lipid to
CC contact APCs, in which receptors for the tumour associated lipids are
CC inactivated or deleted to produce a tumouricidally activated population,

CC and administering APCs to the host), producing a tumouricidal T cell
CC population ex vivo in a mammal) by allowing a tumour associated lipids to
CC contact T cells, in which adaptor proteins, which inhibit T cell
CC activation by tumour associated antigens, are deleted or functionally
CC deactivated to produce a tumouricidal population of T cells, and
CC administering the tumouricidal activated T cells to the host, or
CC allowing a superantigen-lipid raft to contact T cells ex vivo, and
CC administering the tumouricidal activated T cells to the host), treating
CC (M5) cancer in a mammal (by administering a lipid binding molecule which
CC binds immunosuppressive tumour associated lipids in vivo), producing (M6)
CC a tumouricidal T cell population in vivo in a mammal (by allowing a
CC tumour associated antigen to contact immunocytes in which adaptor
CC proteins which inhibit T cell activation by tumour associated antigens
CC are deleted or functionally deactivated) and producing (M7) a
CC tumouricidal T cell population ex vivo in a mammal comprising allowing a
CC superantigen-lipid raft conjugate to contact immunocytes in vivo. The
CC receptors, methods and compositions are useful for treating cancers and
CC tumours. Bacterial superantigens are co-administered or administered as
CC fusion constructs with anti-tumour proteins or moieties. The present
CC sequence encodes an anti-tumour protein which is co-administered with or
CC incorporated into a fusion construct with a superantigen. Note: The
CC sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format from the US patent
CC office website at "seqdata.uspto.gov/sequence.html?docID=20020177551"
CC
XX
SQ Sequence 10422 BP; 3020 A; 2386 C; 2503 G; 2513 T; 0 U; 0 Other;
Query Match 72.0%; Score 144.4; DB 8; Length 10422;
Best Local Similarity 89.5%; Pred. No. 3.1e-39;
Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
QY 9 CTCATCTTTCTAGTCTCTCAAGGCTTCATCTCAAGATCATCTGGTGACACCAA 68
DB 9634 CACTGCTTGAAGAAGTCTCAAGGCTTCATCTCAAGATCATCTGGTGACACCAA 9693
QY 69 GAAGTGAACCTCGAATCTCATGTTCAGGAATRGAAATGTCTTGAAGGCC 128
DB 9694 GAAGTGAACCTCGAATCTCATGTTCAGGAATRGAAATGTCTTGAAGGCC 9753
QY 129 ACACAAAGCATATTTGCTTCTAAAGTAAAGCATGCTGCTCACTG 180
DB 9754 ACACAAAGCATATTTGCTTCTAAAGTAAAGCATGCTGCTCACTG 9805
RESULT 7
ADP43405
ID ADP43405 standard; DNA; 10422 BP.
XX
AC ADP43405;
XX
DT 12-FEB-2004 (first entry)
XX
DE Apolipoprotein polynucleotide seqid 125.
XX
KW receptor; lipid-based tumour associated antigen; cytostatic;
KW antimicrobial; gene therapy; neoplastic disease; tumour; cancer;
KW infectious disease; apolipoprotein; ds.
XX
XX Unidentified.
XX OS
XX US200315713-A1.
XX PN
XX 21-AUG-2003.
XX PD
XX 28-DEC-2000; 2000US-00751708.
XX PF
XX 28-DEC-1999; 99US-0173371P.
XX PR
XX (TERM/) TERMAN D S.
XX PA
XX Terman DS;
XX PI
XX MPI; 2003-787326/74.

DR P-PSDB; ADP43406.
XX
XX New receptor in a mammalian cell that inhibits regular activation by
PT receptors specific for lipid-based tumor associated antigens, useful for
PT treating a neoplastic disease or tumor, and infectious diseases.
XX
XX Example 3; SEQ ID NO 125; 151pp; English.
XX PS
XX The invention describes a receptor in a mammalian cell that inhibits
CC regular activation by receptors specific for lipid-based tumour
CC associated antigen. The receptor has cytostatic and antimicrobial
CC properties and is suitable for use in gene therapy. The receptors,
CC methods and compositions are useful for treating a neoplastic disease or
CC tumour (cancer), and infectious diseases. This sequence represents
CC apolipoprotein polynucleotide, a cell surface moiety, the DNA of which
CC can be transfected into a cell with superantigen DNA to generate
CC antitumour immunity.
CC
XX
SQ Sequence 10422 BP; 3020 A; 2386 C; 2503 G; 2513 T; 0 U; 0 Other;
Query Match 72.0%; Score 144.4; DB 10; Length 10422;
Best Local Similarity 89.5%; Pred. No. 3.1e-39;
Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
QY 9 CTCATCTTTCTAGTCTCTCAAGGCTTCATCTCAAGATCATCTGGTGACACCAA 68
DB 9634 CACTGCTTGAAGAAGTCTCAAGGCTTCATCTCAAGATCATCTGGTGACACCAA 9693
QY 69 GAAGTGAACCTCGAATCTCATGTTCAGGAATRGAAATGTCTTGAAGGCC 128
DB 9694 GAAGTGAACCTCGAATCTCATGTTCAGGAATRGAAATGTCTTGAAGGCC 9753
QY 129 ACACAAAGCATATTTGCTTCTAAAGTAAAGCATGCTGCTCACTG 180
DB 9754 ACACAAAGCATATTTGCTTCTAAAGTAAAGCATGCTGCTCACTG 9805
RESULT 8
AEA03076
ID AEA03076 standard; DNA; 10422 BP.
XX
AC AEA03076;
XX
DT 28-JUL-2005 (first entry)
XX
DE iNOS nucleotide sequence SEQ ID NO:102.
XX
KW tumor; neoplasm; gene therapy; immunotherapy; cytostatic;
KW inducible nitric oxide synthase; gene; ds.
XX
XX Unidentified.
XX OS
XX US2005112141-A1.
XX PN
XX 26-MAY-2005.
XX PD
XX 08-SEP-2004; 2004US-00937758.
XX PF
XX 30-AUG-2000; 2000US-00650884.
XX PR
XX (TERM/) TERMAN D S.
XX PA
XX Terman DS;
XX PI
XX MPI; 2005-394926/40.
XX DR
XX P-PSDB; AEA03077.
XX PT
XX New composition for treating a tumor or neoplastic disease in a subject
PT comprises conjugates comprising superantigen polypeptides or nucleic
XX acids with other molecules that produce a tumoricidal response.
XX
XX Example 3; SEQ ID NO 102; 125pp; English.
XX PS

CC The invention relates to a composition for treating a tumor or neoplastic
CC disease in a subject. Also described: (1) a mammalian cell comprising an
CC exogenous nucleic acid encoding a superantigen expressed in the cell,
CC which cell also produces or expresses all alpha-anomers of
CC monoglycosylceramide or diglycosylceramide, where expression of the
CC superantigen and the mono- or diglycosylceramide is capable of eliciting
CC an antitumor immune response in a mammal into which the cell is
CC introduced; (2) treating a tumor or neoplastic disease in a subject; (3)
CC preparing a population of immunotherapeutic T or natural killer T (NKT)
CC cells useful to treat a tumor or neoplastic disease in a subject; (4) an
CC apoptotic cell preparation or lysate useful for treating a tumor or
CC neoplastic disease in a subject, comprising a cell population that has
CC been transfected with naked DNA encoding a superantigen, and treated to
CC undergo apoptosis or lysis; and (5) a cell that has ingested or been
CC transfected with the above apoptotic preparation or lysate, thus,
CC rendering the cell effective in presenting material expressed from a
CC transfecting nucleic acid or material ingested to the immune system of a
CC mammal to elicit an anti-tumor immune response. The composition and
CC methods are useful for treating tumors or neoplastic diseases. The
CC present sequence represents an INOS nucleotide sequence, which is used in
CC an example from the present invention. Note - The sequence data for this
CC patent is not represented in the printed specification, but was obtained
CC in electronic format directly from the USPTO web site.

XX Sequence 10422 BP; 3020 A; 2386 C; 2503 G; 2513 T; 0 U; 0 Other;

XX Query Match 72.0%; Score 144.4; DB 14; Length 10422;

XX Best Local Similarity 89.5%; Pred. No. 3.1e-39;

XX Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;

XX 9 CTATCTTTTCTAGTCTCTCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCAA 68

XX 9634 CACTGCTTGAAGAAGCTCTCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCAA 9693

XX 69 GAAGTGAACCTCGAATCTCATGTTCAGAAATGAGTGTCTAGGCTGTTCTTGAGACCC 128

XX 9694 GAAGTGAACCTCGAATCTCATGTTCAGAAATGAGTGTCTAGGCTGTTCTTGAGACCC 9753

XX 129 ACACAGACAGATATTTGCTTCTTAAGCTTAAGGCTGCTGCTGCTGCTGCTGCTG 180

XX 9754 ACACAGACAGATATTTGCTTCTTAAGCTTAAGGCTGCTGCTGCTGCTGCTGCTG 9805

XX Db

XX RESULT 9

XX ACC49787

XX ID ACC49787 standard; cDNA; 13869 BP.

XX AC ACC49787;

XX DT 10-JUL-2003 (first entry)

XX Human lipid-associated molecule LIPAM-10 encoding cDNA SEQ ID NO:27.

XX Human; lipid-associated molecule; LIPAM; cytosolic; cerebroprotective;

XX antiarteriosclerotic; anti-HIV; antiallergic; antiparkinsonian; cardiant;

XX anticonvulsant; nootropic; antiinflammatory; antitumor; hepatotropic;

XX antibacterial; virucide; protozoacide; antiparasitic; antipneumic; AIDS;

XX gene therapy; cell proliferative disease; cancer; atherosclerosis; ulcer;

XX autoimmune disease; inflammatory disease; allergy; neurological disorder;

XX stroke; Parkinson's disease; epilepsy; gastrointestinal disorder; obesity;

XX cirrhosis; cardiovascular disorder; myocardial infarction; diabetes;

XX metabolic disorder; developmental disorder; endocrine disorder;

XX pulmonary disorder; infection; lipid metabolism disorder; gene; ss.

PD 27-MAR-2003.

XX 19-SEP-2002; 2002MO-US029980.

XX 21-SEP-2001; 2001US-0324039P.

XX 26-OCT-2001; 2001US-0346197P.

XX 02-NOV-2001; 2001US-0343876P.

XX 30-NOV-2001; 2001US-034211P.

XX 14-DEC-2001; 2001US-0340233P.

XX 18-DEC-2001; 2001US-0342166P.

XX 22-JAN-2002; 2002US-0351262P.

XX 03-MAY-2002; 2002US-037576P.

XX (INCYTE GENOMICS INC.

XX Warren BA, Emerling BM, Lee EA, Chang H, Foreythe IU, Griffin JI,

XX Baughn MR, Chawla NK, Khare R, Reddy R, Lee S, Bulloch SA, Lee SY,

XX Tran UK, Elliott VS, Tang YT, Bhattacharya U, Burrill JD, Blake JJ, Ho A,

XX Zheng W,

XX WPI; 2003-363142/34.

XX P-PsDB; ABR43305.

XX New human lipid-associated molecules (LIPAM) useful for diagnosing,

XX treating and preventing diseases or conditions associated with aberrant

XX LIPAM expression, e.g. cancer, AIDS, atherosclerosis or infections.

XX Claim 5; Page 216-220; 225pp; English.

XX ACC49787 to ACC49794 encode the human lipid-associated molecule proteins

XX given in ABR43296 to ABR43312, designated LIPAM-1 to LIPAM-17 (I). (1)

XX have cytosolic, antiarteriosclerotic, anti-HIV, antiallergic, nootropic,

XX cerebroprotective, antiparkinsonian, anticonvulsant, antiinflammatory,

XX antitumor, hepatotropic, antibacterial, virucide, protozoacide, cardiant,

XX antiparasitic and antipneumic activities, and can be used in gene

XX therapy. The LIPAM polypeptides and polynucleotides are useful in

XX diagnosing, treating and preventing diseases or conditions associated

XX with decreased expression or overexpression of LIPAM, such as cell

XX proliferative diseases (e.g. cancer or atherosclerosis), autoimmune/

XX inflammatory diseases (e.g. AIDS or allergies), neurological disorders

XX (e.g. stroke, Parkinson's disease or epilepsy), gastrointestinal (e.g.

XX ulcer or cirrhosis), cardiovascular (e.g. myocardial infarction),

XX metabolic (e.g. obesity), developmental, endocrine, or pulmonary

XX disorders, infections (e.g. bacterial, viral, parasitic or protozoal),

XX and disorders of lipid metabolism. They are also useful in assessing the

XX effects of exogenous compounds on the expression of nucleic acid and

XX amino acid sequences of LIPAM. The LIPAMs or their fragments are useful

XX in screening compounds for effectiveness as agonist or antagonist of the

XX polypeptides, or in altering the expression of the target polynucleotide

XX and compounds that specifically bind to or modulate the activity of the

XX polypeptide

XX Sequence 13869 BP; 3589 A; 3879 C; 3551 G; 2850 T; 0 U; 0 Other;

XX Query Match 72.0%; Score 144.4; DB 10; Length 13869;

XX Best Local Similarity 89.5%; Pred. No. 3.5e-39;

XX Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;

XX 9 CTATCTTTTCTAGTCTCTCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCAA 68

XX 1150 CACTGCTTGAAGAAGCTCTCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCAA 13209

XX 69 GAAGTGAACCTCGAATCTCATGTTCAGAAATGAGTGTCTAGGCTGTTCTTGAGACCC 128

XX 13210 GAAGTGAACCTCGAATCTCATGTTCAGAAATGAGTGTCTAGGCTGTTCTTGAGACCC 13269

XX Db

XX 129 ACACAGACAGATATTTGCTTCTTAAGCTTAAGGCTGCTGCTGCTGCTGCTGCTG 180

XX 13270 ACACAGACAGATATTTGCTTCTTAAGCTTAAGGCTGCTGCTGCTGCTGCTG 13321

XX Db

XX RESULT 10

XX ACC47280

ID ACC47280 standard; DNA, 13938 BP.
XX
XX ACC47280;
AC
XX
XX 11-AUG-2003 (first entry)
DT
XX
XX Human apolipoprotein(a) encoding DNA.
DE
XX
XX Human apolipoprotein(a) encoding DNA.
KM
XX Apolipoprotein(a); antiarteriosclerotic; cardiant; gene therapy; human;
KM
XX gene; ds.
XX
XX Homo sapiens.
OS
XX
XX
XX Key Location/Qualifiers
XX CDS 46..13692
XX FT /*tag= a
XX FT /product= "apolipoprotein(a)"
XX
XX
XX W02003014307-A2.
XX
XX
XX 20-FEB-2003.
XX
XX
XX 05-AUG-2002; 2002WO-US024920.
XX
XX 07-AUG-2001; 2001US-00923515.
XX
XX (ISIS-) ISIS PHARM INC.
XX
XX Crooke RM, Graham MJ;
XX
XX WPI; 2003-256565/25.
XX
XX P-PSDB; ABR39860.
XX
XX New antisense compound, useful for preparing a composition for treating
PT abnormal lipid or cholesterol metabolism, atherosclerosis or
PT cardiovascular disease.
XX
XX
XX Example 13; Page 92-112, 120pp; English.
XX
XX
XX The invention relates to a new compound, 8-50 nucleobases in length
XX targeted to a nucleic acid molecule encoding human apolipoprotein(a),
XX specifically hybridizes with and inhibits the expression of human
XX apolipoprotein(a). The antisense compounds are useful for preparing a
XX composition for treating abnormal lipid or cholesterol metabolism,
XX atherosclerosis or cardiovascular disease. The present sequence
XX represents a DNA encoding the human apolipoprotein(a) (Genbank accession
XX No. NM_005577)
XX
XX
XX Sequence 13938 BP; 3613 A; 3889 C; 3560 G; 2876 T; 0 U; 0 Other;
SQ
Query Match 72.0%; Score 144.4; DB 8; Length 13938;
Best Local Similarity 89.5%; Pred. No. 3.5e-39;
Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
QY 9 CTCATCTTTTCAGGCTCTCAAGGCTTCATCTCAAGGCTCTGAGGTGACACCA 68
DB 13150 CACTGCTTGAAGAGGCTCTCAAGGCTTCATCTCAAGGCTCTGAGGTGACACCA 13209
QY 69 GAAAGTGAAGCTGGAATCTCATGCTTCAGAAATGAAAGTCTAGAGCTGTTCTTGAAGCC 128
DB 13210 GAAAGTGAAGCTGGAATCTCATGCTTCAGAAATGAAAGTCTAGAGCTGTTCTTGAAGCC 13269
QY 129 ACACAGCAGATATTTGCTTGTCTAAAGCTAAGCAGGTAATCTGCTACCTG 180
DB 13270 ACACAGCAGATATTTGCTTGTCTAAAGCTAAGCAGGCTGCTGCTATCACTG 13321
RESULT 11
ADO33412
ID ADO33412 standard; DNA; 13938 BP.
XX
XX
XX ADO33412;
XX

DT 12-AUG-2004 (first entry)
XX
XX
XX Human apolipoprotein(a) [Lp(a)] DNA - SEQ 860.
DE
XX
XX apolipoprotein B; ApoB; cardiovascular; antiarteriosclerotic;
XX antidiabetic; anorectic; cardiant; vasotropic; hypotensive;
XX anabolic; eating disorder; cytosolic; endocrine; vasotropic;
XX neuroprotective; noctropic; lipid; cholesterol metabolism;
XX hyperlipidaemia; hyperlipidaemia; hypercholesterolaemia;
XX Von Gierke's disease; lipodystrophy; Cushing's syndrome;
XX sexual atelectic dwarfism; hyperthyroidism; hypertension;
XX anorexia nervosa; Werner's syndrome; hepatoma; multiple myeloma; uraemia;
XX impotence; obstructive liver disease; Alzheimer's disease; diabetes;
XX obesity; atherosclerosis; human; ds; apolipoprotein(a); Lp(a).
XX
XX
XX Homo sapiens.
OS
XX
XX W02004044181-A2.
XX
XX
XX 27-MAY-2004.
XX
XX
XX 13-NOV-2003; 2003WO-US036411.
XX
XX
XX 13-NOV-2002; 2002US-0426234P.
XX
XX 15-MAY-2003; 2003WO-US015493.
XX
XX (ISIS-) ISIS PHARM INC.
XX
XX Crooke R, Graham M, Lemonidis-Tardec K, Dobbie KM;
XX
XX WPI; 2004-420321/39.
XX
XX
XX Antisense oligonucleotide compound that inhibits expression of mRNA
XX encoding human apolipoprotein B, useful for treating hyperlipidaemia,
XX diabetes, obesity, von Gierke's disease, lipodystrophies, Cushing's
XX syndrome.
XX
XX
XX Example 57; SEQ ID NO 860; 483pp; English.
XX
XX
XX The invention relates to a novel antisense compound where the compound
XX hybridizes to and inhibits expression of mRNA encoding human
XX apolipoprotein B (ApoB) after 16-24 hours by at least 30% in 80%
XX confluent HepG2 cells in culture at a concentration of 150 nM. The
XX compound of the invention demonstrates cardiovascular,
XX antiarteriosclerotic, antidiabetic, anorectic, cardiant,
XX vasotropic, hypotensive, anabolic, eating disorder-related, cytostatic,
XX endocrine, vasotropic, neuroprotective and noctropic activities and may
XX be useful for inhibiting the expression of apolipoprotein B in cells or
XX tissues in vivo in order to address a condition associated with abnormal
XX lipid or cholesterol metabolism. The compound may be useful for
XX decreasing circulating lipoprotein levels, triglyceride levels,
XX cholesterol levels, lipid levels, fatty acid levels, acute phase
XX reactants and chylomicrons and thus may be utilised during treatment of
XX hyperlipidaemia, hyperlipidaemia, hypercholesterolaemia,
XX cardiovascular disorders, Von Gierke's disease, lipodystrophy, Cushing's
XX syndrome, sexual atelectic dwarfism, hyperthyroidism, hypertension,
XX anorexia nervosa, Werner's syndrome, hepatoma, multiple myeloma, uraemia,
XX impotence, obstructive liver disease, Alzheimer's disease, dementia,
XX diabetes, obesity and atherosclerosis. The current sequence is that of
XX the human apolipoprotein(a) [Lp(a)] DNA - SEQ ID 860 of the invention.
XX
XX
XX Sequence 13938 BP; 3614 A; 3889 C; 3560 G; 2875 T; 0 U; 0 Other;
SQ
Query Match 72.0%; Score 144.4; DB 12; Length 13938;
Best Local Similarity 89.5%; Pred. No. 3.5e-39;
Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;
QY 9 CTCATCTTTTCAGGCTCTCAAGGCTTCATCTCAAGGCTCTGAGGTGACACCA 68
DB 13150 CACTGCTTGAAGAGGCTCTCAAGGCTTCATCTCAAGGCTCTGAGGTGACACCA 13209
QY 69 GAAAGTGAAGCTGGAATCTCATGCTTCAGAAATGAAAGTCTAGAGCTGTTCTTGAAGCC 128
DB 13210 GAAAGTGAAGCTGGAATCTCATGCTTCAGAAATGAAAGTCTAGAGCTGTTCTTGAAGCC 13269
QY 129 ACACAGCAGATATTTGCTTGTCTAAAGCTAAGCAGGTAATCTGCTACCTG 180
DB 13270 ACACAGCAGATATTTGCTTGTCTAAAGCTAAGCAGGCTGCTGCTATCACTG 13321

Db 13310 GAAGTGAACCTCGATCTCATGTTCCAGGAATGAGTGTCTAGGCTGTTCTTGAGCCC 13269

Qy 129 ACACAGAGAGATATTGCTTCTTAAGCTAAGCAGTAAGTCTGCTCAGCTGTG 180
 |||||
 Db 13370 ACACAGAGAGATATTGCTTCTTAAGCTAAGCAGGCGCTGCATCATCTG 13321

RESULT 12

ADQ75139 ID ADQ75139 standard; CDNA; 13938 BP.

AC ADQ75139;

XX 07-OCT-2004 (first entry)

DE Human apolipoprotein A cDNA.

XX cardiovasecular; antiatherosclerotic; antilipemic; vasotropic;
 KM apolipoprotein modulator A; apolipoprotein A; plasminogen;
 KM apolipoprotein A associated disorder; cardiovascular disorder;
 KM atherosclerosis; hypercholesterolaemia; coronary artery disease; human;
 KM gene; 68.

XX Homo sapiens.

XX US2004138164-A1.

PN 15-JUL-2004.

PD 15-JUL-2004.

XX 15-OCT-2003; 2003US-00684440.

PF 15-OCT-2003; 2003US-00684440.

XX 07-AUG-2001; 2001US-00923515.

PR 02-JUN-2003; 2003US-0475402P.

XX (ISIS-) ISIS PHARM INC.

PA Crooke RM, Graham MJ;
 PI WPI; 2004-533399/51.
 DR P-PSDB; ADQ75209.

XX New antisense oligonucleotide compounds, useful for diagnosing,
 PT preventing and/or treating conditions with aberrant expression of
 PT apolipoprotein(a), such as atherosclerosis, hypercholesterolemia and
 PT coronary artery disease.

XX Example 13; SEQ ID NO 4; 54bp; English.

PS The invention describes a new compound (I) comprising 8-80 nucleobases in
 XX length targeted to a nucleic acid molecule encoding apolipoprotein(a),
 CC where the compound is at least 70% complementary to the nucleic acid
 CC molecule encoding apolipoprotein(a), and inhibits the expression of
 CC apolipoprotein(a) mRNA by at least 10%. Also described are: a method of
 CC inhibiting the expression of apolipoprotein(a) in cells or tissues; a
 CC method of treating an animal having a disease or condition associated
 CC with apolipoprotein(a); a method of screening for a modulator of
 CC apolipoprotein(a); a diagnostic method for identifying a disease state,
 CC comprising identifying the presence of apolipoprotein(a) in a sample
 CC using at least one of the primers selected from a fully defined sequence
 CC of 25, 18 or 24 bp (SEQ ID NO: 5, 6 or 7) as given in the specification;
 CC a kit or assay device comprising (I); and a method of inhibiting the
 CC expression of apolipoprotein(a), comprising contacting a biological
 CC system expressing human apolipoprotein(a) with a synthetic antisense
 CC compound, where the synthetic antisense compound comprises 15-30
 CC nucleobases in length and has at least 3 mismatches to human plasminogen.
 CC The methods and compositions of the present invention are useful for the
 CC diagnosis, prevention and/or treatment of diseases or conditions
 CC associated with aberrant expression or activity of apolipoprotein(a),
 CC such as cardiovascular disorder, atherosclerosis, hypercholesterolaemia,
 CC coronary artery disease and/or their combinations. This sequence encodes
 CC human apolipoprotein A.

XX Sequence 13938 BP; 3613 A; 3889 C; 3560 G; 2876 T; 0 U; 0 Other;

Query Match 72.0%; Score 144.4; DB 12; Length 13938;
 Best Local Similarity 89.5%; Pred. No. 3.5e-39;
 Matches 154; Conservative 1; Mismatches 17; Indels 0; Gaps 0;

Qy 9 CTCATCTTTTCTAGTCTCAAGGCTTCATCTCAGAGGTCATCTCGGGTGACACCCA 68
 |||||

Db 13150 CATGCTGAAGAGCTCTAAGGCTTCATCTCAGAGGTCATCTCGGGTGACACCCA 13209

Qy 69 GAAGTGAACCTCGATCTCATGTTCCAGGAATGAGTGTCTAGGCTGTTCTTGAGCCC 128
 |||||

Db 13310 GAAGTGAACCTCGATCTCATGTTCCAGGAATGAGTGTCTAGGCTGTTCTTGAGCCC 13269

Qy 129 ACACAGAGAGATATTGCTTCTTAAGCTAAGCAGTAAGTCTGCTCAGCTGTG 180
 |||||

Db 13370 ACACAGAGAGATATTGCTTCTTAAGCTAAGCAGGCGCTGCATCATCTG 13321

RESULT 13

ADQ38545 ID ADQ38545 standard; DNA; 13938 BP.

XX ADQ38545;

AC ADQ38545;

XX 18-NOV-2004 (first entry)

DE Human SNP containing myocardial infarction-associated gene, SEQ ID 208.

XX Myocardial infarction; detection; single nucleotide polymorphism; SNP;
 KM cardiant; gene therapy; human; gene; de.

XX Homo sapiens.

XX WO2004058052-A2.

XX 15-JUL-2004.

PD 15-JUL-2004.

XX 22-DEC-2003; 2003WO-US040978.

PF 22-DEC-2003; 2003WO-US040978.

XX 20-DEC-2002; 2002US-0434778P.

PR 10-MAR-2003; 2003US-0453135P.

PR 30-APR-2003; 2003US-0466412P.

PR 23-SEP-2003; 2003US-0504955P.

XX (APPL-) APPLERA CORP.

PA Cargill M, Devlin JJ, Iakubova O;
 PI WPI; 2004-533949/51.
 DR P-PSDB; ADQ39373.

XX Identifying an individual who has an altered risk for developing
 PT myocardial infarction by detecting a single nucleotide polymorphism in
 PT the individual's nucleic acids.

XX Claim 7; SEQ ID NO 208; 145bp; English.

PS The invention relates to a novel method for identifying an individual who
 XX has an altered risk for developing myocardial infarction. The method
 CC comprises detecting a single nucleotide polymorphism (SNP) in any one of
 CC the nucleotide sequences given in the specification in the individual's
 CC nucleic acids, where the presence of the SNP is correlated with an
 CC altered risk for myocardial infarction in the individual. The invention
 CC further comprises: an isolated nucleic acid molecule comprising at least
 CC 8 contiguous nucleotides where one of the nucleotides is an SNP given in
 CC the specification or its complement and encoding any one of the amino
 CC acid sequences given in the specification; an isolated polypeptide
 CC comprising an amino acid sequence given in the specification; an antibody
 CC that specifically binds to the polypeptide or its antigen-binding
 CC fragment; an amplified polynucleotide containing an SNP given in the
 CC specification and which is between about 16 and 1000 nucleotides in
 CC length; a kit for detecting an SNP in a nucleic acid, comprising the
 CC polynucleotide, a buffer and an enzyme; a method of detecting an SNP in a

CC nucleic acid molecule; a method of detecting a variant polypeptide; and a
CC method for identifying an agent useful in treating or preventing
CC myocardial infarction. The novel detection method has cardiac activity.
CC The nucleic acids of the invention may be used in gene therapy. The
CC method is useful in identifying an individual who has an increased or
CC decreased risk for developing myocardial infarction and for preparing a
CC composition for treating or preventing myocardial infarction. This
CC polynucleotide sequence represents a human myocardial infarction-
CC associated gene containing one or more SNPs of the invention. Note: This
CC sequence was not shown in the specification. The sequence has come from
CC an electronic sequence listing downloaded from the WIPO website.

XX Sequence 13938 BP; 3611 A; 3880 C; 3553 G; 2873 T; 0 U; 21 Other;

Query Match 72.0%; Score 144.4; DB 13; Length 13938;
Best Local Similarity 90.1%; Pred. No. 3.5e-39;
Matches 155; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

OY 9 CTCATCTTTCTAGGCTCTCAAGGCTTCATCTCTACAGATCATCTGGTGACACCA 68

Db 13150 CACTGCTTGAAGAGTCTCTCAAGGCTTCATCTCTACAGATCATCTGGTGACACCA 13209

OY 69 GAAGTGAACCTCGAATCTCATGTTCAGGAAATRGAAATGCTTACGCTTTCTTGAGGCC 128

Db 13210 GAAGTGAACCTCGAATCTCATGTTCAGGAAATRGAAATGCTTACGCTTTCTTGAGGCC 13269

OY 129 ACACAGAGATATTTGCTTCTTAAGCTTAAGAGCTACTGGCTACCTGTG 180

Db 13270 ACACAGAGATATTTGCTTCTTAAGCTTAAGAGCTACTGGCTACCTGTG 13321

RESULT 14

AAS80436/C
ID AAS80436 standard; cDNA, 2178 BP.

XX AAS80436;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #16240.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KM food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

OS WO200175067-A2.

PN 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US008631.

PR 31-MAR-2000; 2000US-00540217.

PR 23-AUG-2000; 2000US-00649167.

XX (HYSE-) HYSEQ INC.

XX Dymanac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

XX P-PSDB; ABG16249.

PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensic, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity.

XX Claim 1; SEQ ID NO 16240; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
XX sequences. (I) is useful as hybridisation probes, polymerase chain
XX reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
XX and in recombinant production of (II). The polynucleotides are also used

CC in diagnostics as expressed sequence tags for identifying expressed
CC genes. (I) is useful in gene therapy techniques to restore normal
CC activity of (II) or to treat disease states involving (II). (II) is
CC useful for generating antibodies against it, detecting or quantitating a
CC polypeptide in tissue, as molecular weight markers and as a food
CC supplement. (II) and its binding partners are useful in medical imaging
CC of sites expressing (II). (I) and (II) are useful for treating disorders
CC involving aberrant protein expression or biological activity. The
CC polynucleotide and polynucleotide sequences have applications in
CC diagnostics, forensic, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
CC coding sequences of the invention. Note: The sequence data for this
CC patent did not appear in the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences

XX Sequence 2178 BP; 453 A; 532 C; 574 G; 619 T; 0 U; 0 Other;

Query Match 66.2%; Score 132.8; DB 5; Length 2178;
Best Local Similarity 88.3%; Pred. No. 1.7e-35;
Matches 143; Conservative 1; Mismatches 18; Indels 0; Gaps 0;

OY 19 CTAGTCTCTAAGGCTTCTCATCTCAAGATCATCTGGTGACACCAAGAATGAC 78

Db 272 CAAGTCTCTAAGGCTTCTCATCTCAAGATCATCTGGTGACACCAAGAATGAC 213

OY 79 TCGAATCTCATGTTCAGGAAATRGAAATGCTTACGCTTTCTTGAGGCC 138

Db 212 TCGAATCTCATGTTCAGGAAATRGAAATGCTTACGCTTTCTTGAGGCC 153

OY 139 ATATTGCTTCTAAGCTTAAGAGCTACTGGCTACCTGTG 180

Db 152 ATATTGCTTCTAAGCTTAAGAGCTACTGGCTACCTGTG 111

RESULT 15

ACH00783
ID ACH00783 standard; DNA, 744 BP.

XX ACH00783;

DT 12-FEB-2004 (first entry)

DE Human micro-plasminogen gene SEQ ID NO: 64.

KW Recombinant protein production; plasminogen; signal peptide;
KM plasminogen activator identification; protease; wound healing; vulnery;
KM anticoagulant; thrombolytic; cardiac; cerebroprotective; vasotropic;
KM antiinflammatory; ophthalmological; thrombosis; gene; ds.

XX Homo sapiens.

OS WO2003066842-A2.

PN 14-AUG-2003.

XX 06-FEB-2003; 2003WO-DE000341.

PR 06-FEB-2002; 2002EP-00002716.

PR 21-FEB-2002; 2002US-0357809P.

XX (TROM-) TROMMSDORFF ARZNEIMITTEL GMBH & CO KG.

XX Susilo R, Korting HC, Gassen HG, Hils M, Pasternack R;

DR WPI; 2003-697451/66.

XX Recombinant production of plasminogen as a fusion protein comprising a
XX signal peptide that can be removed by proteolysis is useful as an
XX antithrombotic and anticoagulant.

PS Claim 25; Page 68-69; 140pp; German.

XX
CC The present invention relates to a method of producing a recombinant
CC functional plasminogen in microorganisms. The method comprises a sequence
CC encoding at least a functional part of such a plasminogen and a sequence
CC encoding a signal peptide that are fused together, where the two
CC sequences are linked through codons that encode a protease cleavage site
CC to allow removal of the signal peptide. The plasminogen is used to screen
CC for plasminogen activators. Both plasminogen and plasmin, prepared from
CC it by activation, are used for treatment of wounds and the
CC treatment/prevention of thrombotic events, including use in wound
CC dressings, as antithrombotic and anticoagulant agents for preventing or
CC treating a wide variety of conditions, e.g. cardiac infarct, stroke,
CC thrombosis, restenosis, hypoxia, ischemia, vascular inflammation,
CC pulmonary embolism, conjunctivitis (plasminogen type-I deficiency), burns
CC and disseminated intravascular coagulation. The present sequence is a
CC coding sequence used in the exemplification of the invention

XX
SQ Sequence 744 BP; 177 A; 169 C; 207 G; 191 T; 0 U; 0 Other;

Query March 66.1%; Score 132.6; DB 10; Length 744;

Best Local Similarity 85.0%; Pred. No. 1.3e-35;

Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

OY	8	CCTCATCTTTCTAGTCTCAAGGCTTCATCTTACAGGTGATCTGGGTGCACACA	67
Db	174	CCACTGCTTGAGAGAGTCCCAAGGCTTCATCTTACAGGTGATCTGGGTGCACACA	233
OY	68	AGAAAGTGAACCTGCAATCTCATGTTCAAGAAATRGAAGTGTCTAGGCTGTTTGGAGCC	127
Db	234	AGAAAGTGAATCTCGAACCGCATGTTCAAGAAATRGAAGTGTCTAGGCTGTTTGGAGCC	293
OY	128	CACACAGCAGATATGCTGCTTAAGCTTAAGCAGGTACTGGCTCACCTGTG	180
Db	294	CACACGAAAAGATATGCTGCTTAAGCTTAAGCAGGTACTGGCTCACCTGTG	346

Search completed: May 26, 2006, 14:14:02
Job time : 357 secs

RESULT 2
US-09-949-016-15854
; Sequence 15854, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15854
; LENGTH: 55195
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15854

Query Match 92.8%; Score 186.2; DB 3; Length 55195;
Best Local Similarity 95.0%; Pred. No. 9.3e-54;
Matches 191; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

QY 1 TTGACATCTCTCATCTTTTCTAGTCTCTCAAGCCTTCACTCAAGGTCATCTCTGGTG 60
DB 38902 TTGAGCTCTCATCTTTTCTAGTCTCTCAAGCCTTCACTCAAGGTCATCTCTGGTG 38961
QY 61 CACACCAAGAGTGAACCTGATCTCATGTTCAAGAAATGAGTCTAGGCTGTTCT 120
DB 38962 CACACCAAGAGTGAATCTGAAACCGCATGTTCAAGAAATGAGTCTAGGCTGTTCT 39021
QY 121 TGGAGCCCAACACAGAGATATGCTTGTCTAAAGCTAAGAGTCTGCTACCTGTG 180
DB 39022 TGGAGCCCAACAGAAATATGCTTGTCTAAAGCTAAGAGTCTGCTACCTGTG 39081
QY 181 GTTCTTACCCCAAGCTGTGGA 201
DB 39082 GTTCTTACCCCAAGCTGTGGA 39102

RESULT 3
5200340-5
; Patent No. 5200340
; APPLICANT: FOSTER, DONALD C.; MULVIHILL, EILEEN R.; O'HARA,
; PATRICK J.; PINDEL, KURT; YOSHITAKE, SHINJI
; TITLE OF INVENTION: THROMBIN-ACTIVATED TISSUE PLASMINOGEN
; ACTIVATORS
; NUMBER OF SEQUENCES: 34
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/53,412
; FILING DATE: 22-MAY-1987
; SEQ ID NO: 5
; LENGTH: 1724
5200340-5

Query Match 66.1%; Score 132.6; DB 10; Length 1724;
Best Local Similarity 85.0%; Pred. No. 8.9e-36;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTATCTTTTCTAGTCTCTCAAGCCTTCACTCAAGGTCATCTCTGGTGACACCA 67
DB 1086 CCATGCTTGAAGAGTCTCCCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCA 1145
QY 68 AGAAGTGAACCTGATCTCATGTTCAAGAAATGAGTCTAGGCTGTTCTGGAGCC 127
DB 1146 AGAATGAATCTGAAACCGCATGTTCAAGAAATGAGTCTAGGCTGTTCTGGAGCC 1205

QY 128 CACACAGCATATGCTTCTTAAGCTAAGAGGTAAGTCTGCTACCTGTG 180
DB 1206 CACACGAAAGATATGCTTCTTAAGCTAAGAGGTAAGTCTGCTACCTGTG 1258

RESULT 4
US-10-000-489-53
; Sequence 53, Application US/10000489
; Patent No. 6794363
; GENERAL INFORMATION:
; APPLICANT: Benjamin, Stephane
; APPLICANT: Tanaka, Hiroaki
; TITLE OF INVENTION: HUMAN CDNAS AND PROTEINS AND USES THEREOF
; FILE REFERENCE: 91.US.DIV
; CURRENT APPLICATION NUMBER: US/10/000,489
; PRIOR FILING DATE: 2001-11-14
; PRIOR APPLICATION NUMBER: US 09/924,340
; PRIOR FILING DATE: 2001-08-06
; PRIOR APPLICATION NUMBER: PCT/IB01/01715
; PRIOR FILING DATE: 2001-08-06
; PRIOR APPLICATION NUMBER: US 60/305,456
; PRIOR FILING DATE: 2001-07-13
; PRIOR APPLICATION NUMBER: US 60/302,277
; PRIOR FILING DATE: 2001-06-29
; PRIOR APPLICATION NUMBER: US 60/298,698
; PRIOR FILING DATE: 2001-06-15
; PRIOR APPLICATION NUMBER: US 60/293,574
; PRIOR FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 112
; SOFTWARE: JPatent
; SEQ ID NO 53
; LENGTH: 1907
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..1043
; NAME/KEY: CDS
; LOCATION: 1044..1664
; NAME/KEY: 3'UTR
; LOCATION: 1665..1907
; NAME/KEY: polyA_signal
; LOCATION: 1863..1874
; NAME/KEY: polyA_site
; LOCATION: 1892..1907
US-10-000-489-53

Query Match 66.1%; Score 132.6; DB 3; Length 1907;
Best Local Similarity 85.0%; Pred. No. 9.3e-36;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTATCTTTTCTAGTCTCTCAAGCCTTCACTCAAGGTCATCTCTGGTGACACCA 67
DB 1097 CCATGCTTGAAGAGTCTCCCAAGGCTTCACTCAAGGTCATCTCTGGTGACACCA 1156
QY 68 AGAAGTGAACCTGATCTCATGTTCAAGAAATGAGTCTAGGCTGTTCTGGAGCC 127
DB 1157 AGAAGTGAATCTGAAACCGCATGTTCAAGAAATGAGTCTAGGCTGTTCTGGAGCC 1216
QY 128 CACACAGCATATGCTTCTTAAGCTAAGAGGTAAGTCTGCTACCTGTG 180
DB 1217 CACACGAAAGATATGCTTCTTAAGCTAAGAGGTAAGTCTGCTACCTGTG 1269

RESULT 5
US-09-992-095B-53
; Sequence 53, Application US/0992095B
; Patent No. 6989262
; GENERAL INFORMATION:
; APPLICANT: Benjamin, Stephane
; APPLICANT: Tanaka, Hiroaki
; TITLE OF INVENTION: HUMAN CDNAS AND PROTEINS AND USES THEREOF
; FILE REFERENCE: 91.US.DIV

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CURRENT APPLICATION NUMBER: US/09/992,095B
CURRENT FILING DATE: 2003-02-20
PRIOR APPLICATION NUMBER: US 09/924,340
PRIOR FILING DATE: 2001-08-06
PRIOR APPLICATION NUMBER: PCT/IB01/01715
PRIOR FILING DATE: 2001-08-06
PRIOR APPLICATION NUMBER: US 60/305,456
PRIOR FILING DATE: 2001-07-13
PRIOR APPLICATION NUMBER: US 60/302,277
PRIOR FILING DATE: 2001-06-29
PRIOR APPLICATION NUMBER: US 60/298,698
PRIOR FILING DATE: 2001-06-15
PRIOR APPLICATION NUMBER: US 60/293,574
PRIOR FILING DATE: 2001-05-25
NUMBER OF SEQ ID NOS: 112
SOFTWARE: Jpatent
SEQ ID NO 53
LENGTH: 1907
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..1043
FEATURE:
NAME/KEY: CDS
LOCATION: 1044..1664
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 1665..1907
FEATURE:
NAME/KEY: polyA_signal
LOCATION: 1869..1874
FEATURE:
NAME/KEY: polyA_site
LOCATION: 1892..1907
US-09-992-095B-53

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Query Match      66.1%; Score 132.6; DB 4; Length 1907;
Best Local Similarity 85.0%; Pred. No. 9,3e-36;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

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QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCAATCTCTCAAGGCTCTGAGTGCACACCA 67
DB 1097 CCACTGCTTGAGAGAGTCCCAAGGCTTCAATCTCTCAAGGCTCTGAGTGCACACCA 1156
QY 68 AGAAGTGAACCTCGAATTCATGTCAGGAAATGAAAGTGTCTAGGCTGTTTGGAGCC 127
DB 1157 AGAAGTGAATCTCGAAGCGCATGTCAGGAAATGAAAGTGTCTAGGCTGTTTGGAGCC 1216
QY 128 CACACAGCAGATATTGCTTCTTAAGCTAAGCGTAACTGCTCCTCCTGCTG 180
DB 1217 CACACGAAAGATATTGCTTCTTAAGCTAAGCGTAACTGCTCCTGCTGCTGCTG 1269

```

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RESULT 6
US-10-000-986A-53
Sequence 53, Application US/10000986A
Patent No. 7005500
GENERAL INFORMATION:
APPLICANT: Benjamin, Stephanie
APPLICANT: Tanaka, Hiroaki
TITLE OF INVENTION: HUMAN CDNAS AND PROTEINS AND USES THEREOF
FILE REFERENCE: G-0910509D1V
CURRENT APPLICATION NUMBER: US/10/000,986A
CURRENT FILING DATE: 2001-11-14
PRIOR APPLICATION NUMBER: US 09/924,340
PRIOR FILING DATE: 2001-08-06
PRIOR APPLICATION NUMBER: PCT/IB01/01715
PRIOR FILING DATE: 2001-08-06
PRIOR APPLICATION NUMBER: US 60/305,456
PRIOR FILING DATE: 2001-07-13
PRIOR APPLICATION NUMBER: US 60/302,277
PRIOR FILING DATE: 2001-06-29

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PRIOR APPLICATION NUMBER: US 60/298,698
PRIOR FILING DATE: 2001-06-15
PRIOR APPLICATION NUMBER: US 60/293,574
PRIOR FILING DATE: 2001-05-25
NUMBER OF SEQ ID NOS: 228
SOFTWARE: Jpatent
SEQ ID NO 53
LENGTH: 1907
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..1043
FEATURE:
NAME/KEY: CDS
LOCATION: 1044..1664
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 1665..1907
FEATURE:
NAME/KEY: polyA_signal
LOCATION: 1869..1874
FEATURE:
NAME/KEY: polyA_site
LOCATION: 1892..1907
US-10-000-986A-53

```

```

Query Match      66.1%; Score 132.6; DB 5; Length 1907;
Best Local Similarity 85.0%; Pred. No. 9,3e-36;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

```

```

QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCAATCTCTCAAGGCTCTGAGTGCACACCA 67
DB 1097 CCACTGCTTGAGAGAGTCCCAAGGCTTCAATCTCTCAAGGCTCTGAGTGCACACCA 1156
QY 68 AGAAGTGAACCTCGAATTCATGTCAGGAAATGAAAGTGTCTAGGCTGTTTGGAGCC 127
DB 1157 AGAAGTGAATCTCGAAGCGCATGTCAGGAAATGAAAGTGTCTAGGCTGTTTGGAGCC 1216
QY 128 CACACAGCAGATATTGCTTCTTAAGCTAAGCGTAACTGCTCCTCCTGCTGCTG 180
DB 1217 CACACGAAAGATATTGCTTCTTAAGCTAAGCGTAACTGCTCCTCCTGCTGCTGCTG 1269

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RESULT 7
US-07-750-080A-18
Sequence 18, Application US/0750080A
Patent No. 5445953
GENERAL INFORMATION:
APPLICANT: DORNER, F.
APPLICANT: SCHEIFLINGER, F.
APPLICANT: FALKNER, F. G.
TITLE OF INVENTION: DIRECT MOLECULAR CLONING OF A MODIFIED
NUMBER OF SEQUENCES: 42
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 1800 Diagonal Road, Suite 500
CITY: Alexandria
STATE: VA
COUNTRY: USA
ZIP: 22313-0299
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/750,080A
FILING DATE: 19910826
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.

```

REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/106 IMMU
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)833-9300
TELEFAX: (703)833-4109
TELEX: 899149
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 2296 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
IMMEDIATE SOURCE:
CLONE: pN29pc-LPg (Fig. 5.3)
US-07-750-080A-18

Query Match 66.1%; Score 132.6; DB 2; Length 2296;
Best Local Similarity 85.0%; Pred. No. 1e-35;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTCTAGTCTCAAGGCTTCATCTCAAGGTCATCTCGGGTGACACCA 67
DB 1652 CCACTGCTTGAGAGATCCCAAGGCTTCATCTCAAGGTCATCTCGGGTGACACCA 1711

QY 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 127
DB 1712 AGAAGTGAATCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 1771

QY 128 CACACAGCAGATATTGCTTGTCTTAAGCTTAAGCAGAGTACTCGCTACCTGTG 180
DB 1772 CACACGAAAAGATATTGCTTGTCTTAAGCTTAAGCAGAGTACTCGCTACCTGTG 1824

RESULT 8
US-08-651-472-18
Sequence 18, Application US/08651472
Patent No. 6103244
GENERAL INFORMATION:
APPLICANT: DORNER, Friedrich
APPLICANT: SCHEIFLINGER, Friedrich
APPLICANT: FALKNER, Falko Gunter
APPLICANT: FLEIDERER, Michael
TITLE OF INVENTION: DIRECT MOLECULAR CLONING OF CHIMERIC
TITLE OF INVENTION: VIRUSES CONTAINING HUMAN IMMUNODEFICIENCY VIRUS TYPE 1
NUMBER OF SEQUENCES: 95
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/651,472
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/914,738
FILING DATE: 20-JUL-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/750,080
FILING DATE: 26-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/166/IMMU

TELECOMMUNICATION INFORMATION:
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 2296 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Other nucleic acid;
DESCRIPTION: Synthetic DNA oligonucleotide
IMMEDIATE SOURCE:
CLONE: pN29pc-LPg
US-08-651-472-18

Query Match 66.1%; Score 132.6; DB 3; Length 2296;
Best Local Similarity 85.0%; Pred. No. 1e-35;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTCTAGTCTCAAGGCTTCATCTCAAGGTCATCTCGGGTGACACCA 67
DB 1652 CCACTGCTTGAGAGATCCCAAGGCTTCATCTCAAGGTCATCTCGGGTGACACCA 1711

QY 68 AGAAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 127
DB 1712 AGAAGTGAATCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 1771

QY 128 CACACAGCAGATATTGCTTGTCTTAAGCTTAAGCAGAGTACTCGCTACCTGTG 180
DB 1772 CACACGAAAAGATATTGCTTGTCTTAAGCTTAAGCAGAGTACTCGCTACCTGTG 1824

RESULT 9
US-08-358-928-18
Sequence 18, Application US/08358928
Patent No. 6265183
GENERAL INFORMATION:
APPLICANT: DORNER, Friedrich
APPLICANT: SCHEIFLINGER, Friedrich
APPLICANT: FALKNER, Falko Gunter
APPLICANT: FLEIDERER, Michael
TITLE OF INVENTION: DIRECT MOLECULAR CLONING OF CHIMERIC
TITLE OF INVENTION: VIRUSES CONTAINING HUMAN IMMUNODEFICIENCY VIRUS TYPE 1
NUMBER OF SEQUENCES: 95
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington
STATE: D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/358,928
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/914,738
FILING DATE: 20-JUL-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/750,080
FILING DATE: 26-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/166/IMMU
TELECOMMUNICATION INFORMATION:

TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 18:
SEQUENCE CHARACTERISTICS:
LENGTH: 2296 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Other nucleic acid;
DESCRIPTION: Synthetic DNA oligonucleotide
IMMEDIATE SOURCE:
CLONE: pN29pt-Lp9
US-08-358-928-18

Query Match 66.1%; Score 132.6; DB 3; Length 2296;
Best Local Similarity 85.0%; Pred. No. 1e-35;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 67
DB 1652 CCACGTGTTGAGAGAGTCCCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 1711
QY 68 AGAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 127
DB 1712 AGAAGTGAATCTCGAATCCGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 1771
QY 128 CACACAGCAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 180
DB 1772 CACACGAAAAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 1824

RESULT 10
US-09-192-012-4
Sequence 4, Application US/09192012A
Patent No. 6475784
GENERAL INFORMATION:
APPLICANT: Packoff, Jackie
APPLICANT: Megabios Corporation
APPLICANT: Pfizer, Inc.
TITLE OF INVENTION: Inhibition of Angiogenesis by Delivery of Nucleic Acids
TITLE OF INVENTION: Encoding Anti-Angiogenesis Polypeptides
FILE REFERENCE: 018484-000110US
CURRENT APPLICATION NUMBER: US/09/192,012A
CURRENT FILING DATE: 1998-11-13
EARLIER APPLICATION NUMBER: US 60/066,020
EARLIER FILING DATE: 1997-11-14
NUMBER OF SEQ ID NOS: 9
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 4
LENGTH: 2430
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (1) -(2430)
OTHER INFORMATION: human plasminogen
US-09-192-012-4

Query Match 66.1%; Score 132.6; DB 3; Length 2430;
Best Local Similarity 85.0%; Pred. No. 1e-35;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 67
DB 1863 CCACGTGTTGAGAGAGTCCCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 1922
QY 68 AGAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 127
DB 1923 AGAAGTGAATCTCGAATCCGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 1982
QY 128 CACACAGCAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 180

DB 1983 CACACGAAAAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 2035

RESULT 11
US-08-643-219-12
Sequence 12, Application US/08643219
Patent No. 5801146
GENERAL INFORMATION:
APPLICANT: Davidson, Donald J.
TITLE OF INVENTION: NOVEL ANTIANGIOGENIC PEPTIDES
TITLE OF INVENTION: AND METHODS FOR INHIBITING ANGIOGENESIS
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/643,219
FILING DATE: 06-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Casuto, Dianne
REGISTRATION NUMBER: 40,943
REFERENCE/DOCKET NUMBER: 5940.US.P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847-938-3137
TELEFAX: 847-938-2623
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 2497 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-08-643-219-12

Query Match 66.1%; Score 132.6; DB 2; Length 2497;
Best Local Similarity 85.0%; Pred. No. 1e-35;
Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;

QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 67
DB 1912 CCACGTGTTGAGAGAGTCCCAAGGCTTCATCTCAAGGTCACTCTGGGTGCACACCA 1971
QY 68 AGAAGTGAACCTCGAATCTCATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 127
DB 1972 AGAAGTGAATCTCGAATCCGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCTTGAGCC 2031
QY 128 CACACAGCAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 180
DB 2032 CACACGAAAAGATATTTGCTTGTCTTAAGCTTAAGAGGTACTCGCTCACTGTG 2084

RESULT 12
US-09-131-995-12
Sequence 12, Application US/09131995
Patent No. 5972896
GENERAL INFORMATION:
APPLICANT: Davidson, Donald J.
TITLE OF INVENTION: NOVEL ANTIANGIOGENIC PEPTIDES
TITLE OF INVENTION: AND METHODS FOR INHIBITING ANGIOGENESIS

NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FaetsEQ Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/131,995
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/832,087
FILING DATE: 03-APR-1997
APPLICATION NUMBER: 08/643,219
FILING DATE: 06-MAY-1996
ATTORNEY/AGENT INFORMATION:
NAME: Casuto, Dianne
REGISTRATION NUMBER: 40,943
REFERENCE/DOCKET NUMBER: 5940.US.P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847-938-3137
TELEFAX: 847-938-2623
TELEX:
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 2497 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-09-131-995-12

Query Match 66.1%; Score 132.6; DB 2; Length 2497;
Best Local Similarity 85.0%; Pred. No. 1e-35; Indels 0; Gaps 0;
Matches 147; Conservative 1; Mismatches 25;
Qy 8 CCTCATCTTTCTAGTCTCAAGGCTTCACTCTACAGGCTTCTGAGTGCACACCA 67
Db 1912 CCACCTGCTTGAAGAGAGTCCCAAGGCTTCTATCTTACAGGTATCTGGGTGCACACCA 1971
Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAGAAATRGAAAGTGTCTAGGCTGTTCTTGAAGCC 127
Db 1972 AGAAGTGAATCTCGAAGCGCATGTTCAAGAAATAGAGTGTCTAGGCTGTTCTTGAAGCC 2031
Qy 128 CACACAGCAGATATTGCTTGTCTTAAAGCTAAGCAGGTAAGTCTGCTCAGCTGTG 180
Db 2032 CACACGAAAGATATTGCTTGTCTTAAAGCTAAGCAGGTAAGTCTGCTCAGCTGTG 2084

RESULT 13
US-08-832-087B-12
Sequence 12, Application US/08832087B
Patent No. 5981484
GENERAL INFORMATION:
APPLICANT: Davidson, Donald J.
TITLE OF INVENTION: NOVEL ANTIANGIOGENIC PEPTIDES
TITLE OF INVENTION: AND METHODS FOR INHIBITING ANGIOGENESIS
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064
COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FaetsEQ Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/832,087B
FILING DATE: 03-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/643,219
FILING DATE: 06-MAY-1996
ATTORNEY/AGENT INFORMATION:
NAME: Casuto, Dianne
REGISTRATION NUMBER: 40,943
REFERENCE/DOCKET NUMBER: 5940.US.P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847-938-3137
TELEFAX: 847-938-2623
TELEX:
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 2497 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-08-832-087B-12

Query Match 66.1%; Score 132.6; DB 2; Length 2497;
Best Local Similarity 85.0%; Pred. No. 1e-35; Indels 0; Gaps 0;
Matches 147; Conservative 1; Mismatches 25;
Qy 8 CCTCATCTTTCTAGTCTCAAGGCTTCACTCTACAGGCTTCTGAGTGCACACCA 67
Db 1912 CCACCTGCTTGAAGAGTCCCAAGGCTTCTATCTTACAGGTATCTGGGTGCACACCA 1971
Qy 68 AGAAGTGAACCTCGAATCTCATGTTCAGAAATRGAAAGTGTCTAGGCTGTTCTTGAAGCC 127
Db 1972 AGAAGTGAATCTCGAAGCGCATGTTCAAGAAATAGAGTGTCTAGGCTGTTCTTGAAGCC 2031
Qy 128 CACACAGCAGATATTGCTTGTCTTAAAGCTAAGCAGGTAAGTCTGCTCAGCTGTG 180
Db 2032 CACACGAAAGATATTGCTTGTCTTAAAGCTAAGCAGGTAAGTCTGCTCAGCTGTG 2084

RESULT 14
US-08-851-350-12
Sequence 12, Application US/08851350
Patent No. 6057122
GENERAL INFORMATION:
APPLICANT: Abbott Laboratories
TITLE OF INVENTION: NOVEL ANTIANGIOGENIC PEPTIDES,
TITLE OF INVENTION: POLYNUCLEOTIDES ENCODING SAME AND METHODS
TITLE OF INVENTION: FOR INHIBITING ANGIOGENESIS
NUMBER OF SEQUENCES: 38
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FaetsEQ Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/851,350
FILING DATE: 05-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER:

FILING DATE:
 ATTORNEY/AGENT INFORMATION:
 NAME: Casuto, Dianne
 REGISTRATION NUMBER: 40,943
 REFERENCE/DOCKET NUMBER: 5940 US, P2
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 847-938-2137
 TELEFAX: 847-938-2623
 TELEX:
 INFORMATION FOR SEQ ID NO: 12:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 2497 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 US-08-851-350-12

Query Match 66.1%; Score 132.6; DB 3; Length 2497;
 Best Local Similarity 85.0%; Pred. No. 1e-35;
 Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;
 QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCAATCTTCAAGGCTATCTGAGGCTGACACCA 67
 DB 1912 CCACTGCTTGAGAGAGTCCCAAGGCTTCAATCTTCAAGGCTATCTGAGGCTGACACCA 1971
 QY 68 AGAAGTGAACCTCGAATCTCATGTTTCAGGAATGAAAGTCTAGGCTGTTCTTGAGACC 127
 DB 1972 AGAAGTGAATCTCGAATCTCATGTTTCAGGAATGAAAGTCTAGGCTGTTCTTGAGACC 2031
 QY 128 CACACAGCAGATATGCTGCTTAAAGCTAAGCAGGTAAGTCTGCTACCTGTG 180
 DB 2032 CACACGAAAAGATATGCTGCTTAAAGCTAAGCAGGTAAGTCTGCTACCTGTG 2084

RESULT 15
 US-09-132-154-12
 Sequence 12, Application US/09132154
 Patent No. 6251867
 GENERAL INFORMATION:
 APPLICANT: Davidson, Donald J.
 TITLE OF INVENTION: NOVEL ANTIANGIOGENIC PEPTIDES
 TITLE OF INVENTION: AND METHODS FOR INHIBITING ANGIOGENESIS
 NUMBER OF SEQUENCES: 14
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Abbott Laboratories
 STREET: 100 Abbott Park Road
 CITY: Abbott Park
 STATE: IL
 COUNTRY: USA
 ZIP: 60064
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: Pasteo Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/132,154
 FILING DATE:
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US/08/832,087
 FILING DATE: 03-APR-1997
 APPLICATION NUMBER: 08/643,219
 FILING DATE: 06-MAY-1996
 ATTORNEY/AGENT INFORMATION:
 NAME: Casuto, Dianne
 REGISTRATION NUMBER: 40,943
 REFERENCE/DOCKET NUMBER: 5940 US, P1
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 847-938-2137
 TELEFAX: 847-938-2623
 TELEX:
 INFORMATION FOR SEQ ID NO: 12:

SEQUENCE CHARACTERISTICS:
 LENGTH: 2497 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: double
 TOPOLOGY: linear
 MOLECULE TYPE: cDNA
 US-09-132-154-12

Query Match 66.1%; Score 132.6; DB 3; Length 2497;
 Best Local Similarity 85.0%; Pred. No. 1e-35;
 Matches 147; Conservative 1; Mismatches 25; Indels 0; Gaps 0;
 QY 8 CCTCATCTTTCTAGGCTCTCAAGGCTTCAATCTTCAAGGCTATCTGAGGCTGACACCA 67
 DB 1912 CCACTGCTTGAGAGAGTCCCAAGGCTTCAATCTTCAAGGCTATCTGAGGCTGACACCA 1971
 QY 68 AGAAGTGAACCTCGAATCTCATGTTTCAGGAATGAAAGTCTAGGCTGTTCTTGAGACC 127
 DB 1972 AGAAGTGAATCTCGAATCTCATGTTTCAGGAATGAAAGTCTAGGCTGTTCTTGAGACC 2031
 QY 128 CACACAGCAGATATGCTGCTTAAAGCTAAGCAGGTAAGTCTGCTACCTGTG 180
 DB 2032 CACACGAAAAGATATGCTGCTTAAAGCTAAGCAGGTAAGTCTGCTACCTGTG 2084

Search completed: May 26, 2006, 13:54:15
 Job time : 96 secs


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RESULT 2
US-10-995-561-25501
; Sequence 25501, Application US/10995561
; Publication No. US20050272054A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; TITLE OF INVENTION: DETECTION AND USES THEREOF
; FILE REFERENCE: CL001559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 25501
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-995-561-25501
Query Match 100.0%; Score 200.6; DB 10; Length 201;
Best Local Similarity 100.0%; Pred. No. 4,5e-61;
Matches 201; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 60
DB 1 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 60

QY 61 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 120
DB 61 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 120

QY 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180
DB 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180

QY 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180
DB 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180

QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 181 GTCTTACCCCAAGCTGTGTA 201

RESULT 3
US-11-124-368A-8354/C
; Sequence 8354, Application US/11124368A
; Publication No. US20050287559A1
; GENERAL INFORMATION:
; APPLICANT: Michele Cargill
; APPLICANT: James J. Devlin
; TITLE OF INVENTION: Genetic Polymorphisms Associated with
; TITLE OF INVENTION: Vascular Diseases, Methods of Detection and Uses Thereof
; FILE REFERENCE: CL001524
; CURRENT APPLICATION NUMBER: US/11/124,368A
; CURRENT FILING DATE: 2005-05-09
; PRIOR APPLICATION NUMBER: US 60/566,845
; PRIOR FILING DATE: 2004-05-07
; PRIOR APPLICATION NUMBER: US 60/625,936
; PRIOR FILING DATE: 2004-11-09
; NUMBER OF SEQ ID NOS: 21112
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 8354
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-124-368A-8354
Query Match 100.0%; Score 200.6; DB 15; Length 201;
Best Local Similarity 100.0%; Pred. No. 4,5e-61;
Matches 201; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 60
DB 201 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 142
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QY 61 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 120
DB 141 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 82

QY 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180
DB 81 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 22

QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 21 GTCTTACCCCAAGCTGTGTA 1

RESULT 4
US-10-741-600-17618
; Sequence 17618, Application US/10741600
; Publication No. US20050026169A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: MYOCARDIAL INFARCTION, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001499
; CURRENT APPLICATION NUMBER: US/10/741,600
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 73997
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17618
; LENGTH: 93112
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(93112)
; OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables)
US-10-741-600-17618
Query Match 100.0%; Score 200.6; DB 9; Length 93112;
Best Local Similarity 100.0%; Pred. No. 5,6e-60;
Matches 201; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 60
DB 79990 TTGACATCTCATCTTTTCTAGGTCCTCAAGGCTTCATCTCAAGGTCACTCTGGGTG 79049

QY 61 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 120
DB 79050 CACACCAAGAAGTGAAGTGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 79109

QY 121 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 180
DB 79110 TGGAGCCCAACACAGAGATATTCCTTGTCTAAGAGTAAAGTAACTGCTCACTGTG 79169

QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 79170 GTCTTACCCCAAGCTGTGTA 79190

RESULT 5
US-10-995-561-13234
; Sequence 13234, Application US/10995561
; Publication No. US20050272054A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; TITLE OF INVENTION: DETECTION AND USES THEREOF
; FILE REFERENCE: CL001559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13234
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/
/   LENGTH: 93112
/   TYPE: DNA
/   ORGANISM: Homo sapiens
/   FEATURE:
/   NAME/KEY: misc_feature
/   LOCATION: (1)...(93112)
/   OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
US-10-995-561-113234
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Query Match      100.0%; Score 200.6; DB 10; Length 93112;
Best Local Similarity 100.0%; Pred. No. 5.6e-60;
Matches 201; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 1 TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 60
DB TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 79049
QY 61 CACACCAAGAGTGAACCTCGAATCTGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 120
DB CACACCAAGAGTGAACCTCGAATCTGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 79109
QY 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 180
DB 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 79169
QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 181 GTCTTACCCCAAGCTGTGTA 79190
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RESULT 6

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US-09-925-065A-869309/c
/ Sequence 869309, Application US/09925065A
/ Publication No. US20040181048A1
/ GENERAL INFORMATION:
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single
/ TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.135
/ CURRENT APPLICATION NUMBER: US/09/925,065A
/ CURRENT FILING DATE: 2001-08-08
/ PRIOR APPLICATION NUMBER: US 60/243,096
/ PRIOR FILING DATE: 2000-10-24
/ PRIOR APPLICATION NUMBER: US 60/252,147
/ PRIOR FILING DATE: 2000-11-20
/ PRIOR APPLICATION NUMBER: US 60/250,092
/ PRIOR FILING DATE: 2000-11-30
/ PRIOR APPLICATION NUMBER: US 60/261,766
/ PRIOR FILING DATE: 2001-01-16
/ PRIOR APPLICATION NUMBER: US 60/289,846
/ PRIOR FILING DATE: 2001-05-09
/ NUMBER OF SEQ ID NOS: 957086
/ SOFTWARE: FastSeq for Windows Version 4.0
/ SEQ ID NO 869309
/ LENGTH: 614
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-925-065A-869309
```

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Query Match      93.4%; Score 187.4; DB 4; Length 614;
Best Local Similarity 95.0%; Pred. No. 3.8e-56;
Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;
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```
QY 1 TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 60
DB 406 TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 347
QY 61 CACACCAAGAGTGAACCTCGAATCTGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 120
DB 346 CACACCAAGAGTGAATCTCGAAGCCGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 287
QY 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 180
DB 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 180
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DB 286 TGAAGCCCAACAAGAAATATTGCTTCTTAAGTAAAGCAGTACTCGTTCACCTGTG 227
QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 226 GTCTTACCCCAAGCTGTGTA 206
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RESULT 7

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US-09-925-065A-886262/c
/ Sequence 886262, Application US/09925065A
/ Publication No. US20040181048A1
/ GENERAL INFORMATION:
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single
/ TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.135
/ CURRENT APPLICATION NUMBER: US/09/925,065A
/ CURRENT FILING DATE: 2001-08-08
/ PRIOR APPLICATION NUMBER: US 60/243,096
/ PRIOR FILING DATE: 2000-10-24
/ PRIOR APPLICATION NUMBER: US 60/252,147
/ PRIOR FILING DATE: 2000-11-20
/ PRIOR APPLICATION NUMBER: US 60/250,092
/ PRIOR FILING DATE: 2000-11-30
/ PRIOR APPLICATION NUMBER: US 60/261,766
/ PRIOR FILING DATE: 2001-01-16
/ PRIOR APPLICATION NUMBER: US 60/289,846
/ PRIOR FILING DATE: 2001-05-09
/ NUMBER OF SEQ ID NOS: 957086
/ SOFTWARE: FastSeq for Windows Version 4.0
/ SEQ ID NO 886262
/ LENGTH: 614
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-925-065A-886262
```

```
Query Match      93.4%; Score 187.4; DB 4; Length 614;
Best Local Similarity 95.0%; Pred. No. 3.8e-56;
Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;
```

```
QY 1 TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 60
DB 406 TTGACATCCCTCATCTTTCTTAGGTCCTCAAGGCTTCTCAAGAGTCTATCCTGGGTG 347
QY 61 CACACCAAGAGTGAACCTCGAATCTGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 120
DB 346 CACACCAAGAGTGAATCTCGAAGCCGATGTTTCAGAAATGAAAGTGTCTAGGCTGTTCT 287
QY 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 180
DB 121 TGAAGCCCAACAAGAGATATTGCTTGTAAAGCTTAAGCAGTACTCGCTCACTGTG 227
QY 181 GTCTTACCCCAAGCTGTGTA 201
DB 181 GTCTTACCCCAAGCTGTGTA 206
```

RESULT 8

```
US-09-925-065A-869309/c
/ Sequence 869309, Application US/09925065A
/ Publication No. US20050228172A9
/ GENERAL INFORMATION:
/ APPLICANT: Wang, David G.
/ TITLE OF INVENTION: Identification and Mapping of Single
/ TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
/ FILE REFERENCE: 108827.135
/ CURRENT APPLICATION NUMBER: US/09/925,065A
/ CURRENT FILING DATE: 2001-08-08
/ PRIOR APPLICATION NUMBER: US 60/243,096
/ PRIOR FILING DATE: 2000-10-24
/ PRIOR APPLICATION NUMBER: US 60/252,147
/ PRIOR FILING DATE: 2000-11-20
/ PRIOR APPLICATION NUMBER: US 60/250,092
```

PRIOR FILING DATE: 2000-11-30
PRIOR APPLICATION NUMBER: US 60/261,766
PRIOR FILING DATE: 2001-01-16
PRIOR APPLICATION NUMBER: US 60/289,846
PRIOR FILING DATE: 2001-05-09
NUMBER OF SEQ ID NOS: 957086
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 869309
LENGTH: 614
TYPE: DNA
ORGANISM: Homo sapiens
US-09-925-065A-869309

Query Match 93.4%; Score 187.4; DB 5; Length 614;
Best Local Similarity 95.0%; Pred. No. 3.8e-56;
Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 60
DB 406 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 347
QY 61 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 120
DB 346 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 287
QY 121 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 180
DB 286 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 227
QY 181 GTCTTACCCCAACGCTGTGTA 201
DB 226 GTCTTACCCCAACGCTGTGTA 206

RESULT 9

US-09-925-065A-886262/c
Sequence 886262, Application US/09925065A
Publication No. US2005028172A9
GENERAL INFORMATION:
APPLICANT: Wang, David G.
TITLE OF INVENTION: Identification and Mapping of Single
FILE REFERENCE: 108827.135
CURRENT APPLICATION NUMBER: US/09/925,065A
CURRENT FILING DATE: 2001-08-08
PRIOR APPLICATION NUMBER: US 60/243,096
PRIOR FILING DATE: 2000-10-24
PRIOR APPLICATION NUMBER: US 60/252,147
PRIOR FILING DATE: 2000-11-20
PRIOR APPLICATION NUMBER: US 60/250,092
PRIOR FILING DATE: 2000-11-30
PRIOR APPLICATION NUMBER: US 60/261,766
PRIOR FILING DATE: 2001-01-16
PRIOR APPLICATION NUMBER: US 60/289,846
PRIOR FILING DATE: 2001-05-09
NUMBER OF SEQ ID NOS: 957086
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 886262
LENGTH: 614
TYPE: DNA
ORGANISM: Homo sapiens
US-09-925-065A-886262

Query Match 93.4%; Score 187.4; DB 5; Length 614;
Best Local Similarity 95.0%; Pred. No. 3.8e-56;
Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 60
DB 406 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 347
QY 61 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 120

DB 346 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 287
QY 121 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 180
DB 286 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 227
QY 181 GTCTTACCCCAACGCTGTGTA 201
DB 226 GTCTTACCCCAACGCTGTGTA 206

RESULT 10

US-10-741-601-5650
Sequence 5650, Application US/10741601
Publication No. US20040166519A1
GENERAL INFORMATION:
APPLICANT: CARGILL, Michele et al.
TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
FILE REFERENCE: CL001500
CURRENT APPLICATION NUMBER: US/10/741,601
CURRENT FILING DATE: 2003-12-22
NUMBER OF SEQ ID NOS: 26415
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 5650
LENGTH: 63693
TYPE: DNA
ORGANISM: Homo sapiens
US-10-741-601-5650

Query Match 93.4%; Score 187.4; DB 8; Length 63693;
Best Local Similarity 95.0%; Pred. No. 2.6e-55;
Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 60
DB 42930 TTGACATCTCATCTTTTCTAGTCTTCAGGCGCTTCACTTCAAGGTCATCTGGGTG 42989
QY 61 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 120
DB 42990 CACACCAAGAAGTGAAGTCAATCTCATCTTCAAGGTAATGAAGTCTGAGGCTGTTCT 43049
QY 121 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 180
DB 43050 TGGAGCCCAACAAGATATTTGCTTGTAAAGCTAAGGAGTACGCTCACTGTG 43109
QY 181 GTCTTACCCCAACGCTGTGTA 201
DB 43110 GTCTTACCCCAACGCTGTGTA 43130

RESULT 11

US-10-995-561-13269
Sequence 13269, Application US/10995561
Publication No. US20050272054A1
GENERAL INFORMATION:
APPLICANT: CARGILL, Michele et al.
TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
FILE REFERENCE: CL001559
CURRENT APPLICATION NUMBER: US/10/995,561
CURRENT FILING DATE: 2004-11-24
NUMBER OF SEQ ID NOS: 85702
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 13269
LENGTH: 63693
TYPE: DNA
ORGANISM: Homo sapiens
US-10-995-561-13269

Query Match 93.4%; Score 187.4; DB 10; Length 63693;
Best Local Similarity 95.0%; Pred. No. 2.6e-55;

Matches 191; Conservative 2; Mismatches 8; Indels 0; Gaps 0;

```
QY 1 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 60
    |||
DB 42930 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 42989
QY 61 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 120
    |||
DB 42990 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 43049
QY 121 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 180
    |||
DB 43050 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 43109
QY 181 GTCTTACCCCAACGCTGTGTA 201
    |||
DB 43110 GTCTTACCCCAACGCTGTGTA 43130
```

RESULT 12

```
US-11-108-459-9
; Sequence 9, Application US/11108459
; Publication No. US20060051780A1
; GENERAL INFORMATION:
; APPLICANT: Schwartz, David A.
; APPLICANT: Pelitz, Gary
; TITLE OF INVENTION: Polymorphic Plasminogen Genes and Uses Thereof
; FILE REFERENCE: DIXE-09785
; CURRENT APPLICATION NUMBER: US/11/108,459
; CURRENT FILING DATE: 2005-04-18
; NUMBER OF SEQ ID NOS: 9
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 9
; LENGTH: 52280
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-108-459-9
```

Query Match 92.8%; Score 186.2; DB 16; Length 52280;

Best Local Similarity 95.0%; Pred. No. 6.4e-55; Indels 0; Gaps 0;

```
Matches 191; Conservative 1; Mismatches 9; Indels 0; Gaps 0;
QY 1 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 60
    |||
DB 37409 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 37468
QY 61 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 120
    |||
DB 37469 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 37528
QY 121 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 180
    |||
DB 37529 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 37588
QY 181 GTCTTACCCCAACGCTGTGTA 201
    |||
DB 37589 GTCTTACCCCAACGCTGTGTA 37609
```

RESULT 13

```
US-10-741-601-14344
; Sequence 14344, Application US/10741601
; Publication No. US2004016519A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: STENOSIS, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01500
; CURRENT APPLICATION NUMBER: US/10/741,601
; CURRENT FILING DATE: 2003-12-22
; NUMBER OF SEQ ID NOS: 26415
; SOFTWARE: PatSeq for Windows Version 4.0
```

```
; SEQ ID NO 14344
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-741-601-14344
```

Query Match 86.1%; Score 172.8; DB 8; Length 201;

Best Local Similarity 95.2%; Pred. No. 4.2e-51; Indels 0; Gaps 0;

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Matches 177; Conservative 1; Mismatches 8; Indels 0; Gaps 0;
QY 1 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 60
    |||
DB 16 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 75
QY 61 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 120
    |||
DB 76 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 135
QY 121 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 180
    |||
DB 136 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 195
QY 181 GTCTTC 186
    |||
DB 196 GTCTTC 201
```

RESULT 14

```
US-10-995-561-34498
; Sequence 34498, Application US/10995561
; Publication No. US20050272054A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; TITLE OF INVENTION: DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: PatSeq for Windows Version 4.0
; SEQ ID NO 34498
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-995-561-34498
```

Query Match 86.1%; Score 172.8; DB 10; Length 201;

Best Local Similarity 95.2%; Pred. No. 4.2e-51; Indels 0; Gaps 0;

```
Matches 177; Conservative 1; Mismatches 8; Indels 0; Gaps 0;
QY 1 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 60
    |||
DB 16 TTGACATCCCTCATCTTTCTAGGCTCCCAAGGCTTCATCTCAAGGTCATCTGGGTG 75
QY 61 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 120
    |||
DB 76 CACACCAAGAGTGAATCTGAAATCTCATGTTCAAGAAATGAAAGTGTAGGCTGTCT 135
QY 121 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 180
    |||
DB 136 TGAAGCCCAACACAGCAGATATGCTTGTCTAAAGCTAAGAGTATCTGCTACCTGTG 195
QY 181 GTCTTC 186
    |||
DB 196 GTCTTC 201
```

RESULT 15

```
US-11-124-368A-2896
; Sequence 2896, Application US/11124368A
; Publication No. US20050287559A1
; GENERAL INFORMATION:
```

```

1 APPLICANT: Michael Cargill
2 APPLICANT: James J. Devlin
3 APPLICANT: May Luke
4 TITLE OF INVENTION: Genetic Polymorphisms Associated with
5 TITLE OF INVENTION: Vascular Diseases, Methods of Detection and Uses Thereof
6 FILE REFERENCE: CU001524
7 CURRENT APPLICATION NUMBER: US/11/124,368A
8 CURRENT FILING DATE: 2005-05-09
9 PRIOR APPLICATION NUMBER: US 60/566,845
10 PRIOR FILING DATE: 2004-05-07
11 PRIOR APPLICATION NUMBER: US 60/625,936
12 PRIOR FILING DATE: 2004-11-09
13 NUMBER OF SEQ ID NOS: 2112
14 SOFTWARE: FASTSEQ for Windows Version 4.0
15 SEQ ID NO 2896
16 LENGTH: 91561
17 TYPE: DNA
18 ORGANISM: Homo sapiens
19 FEATURE:
20 NAME/KEY: misc_feature
21 LOCATION: 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17,
22 LOCATION: 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31,
23 LOCATION: 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45,
24 LOCATION: 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59,
25 LOCATION: 60
26 OTHER INFORMATION: n = A,T,C or G
27 FEATURE:
28 NAME/KEY: misc_feature
29 LOCATION: 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75,
30 LOCATION: 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89,
31 LOCATION: 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103,
32 LOCATION: 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114
33 OTHER INFORMATION: n = A,T,C or G
34 FEATURE:
35 NAME/KEY: misc_feature
36 LOCATION: 115, 116, 117, 118, 119, 120, 121, 122, 123, 124, 125, 126,
37 LOCATION: 127, 128, 129, 130, 131, 132, 133, 134, 135, 136, 137, 138,
38 LOCATION: 139, 140, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150,
39 LOCATION: 151, 152, 153, 154, 155, 156, 157, 158, 159, 160, 161
40 OTHER INFORMATION: n = A,T,C or G
41 FEATURE:
42 NAME/KEY: misc_feature
43 LOCATION: 162, 163, 164, 165, 166, 167, 168, 169, 170, 171, 172, 173,
44 LOCATION: 174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184, 185,
45 LOCATION: 186, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196, 197,
46 LOCATION: 198, 199, 200, 201, 202, 203, 204, 205, 206, 207, 208
47 OTHER INFORMATION: n = A,T,C or G
48 FEATURE:
49 NAME/KEY: misc_feature
50 LOCATION: 209, 210, 211, 212, 213, 214, 215, 216, 217, 218, 219, 220,
51 LOCATION: 221, 222, 223, 224, 225, 226, 227, 228, 229, 230, 231, 232,
52 LOCATION: 233, 234, 235, 236, 237, 238, 239, 240, 241, 242, 243, 244,
53 LOCATION: 245, 246, 247, 248, 249, 250, 251, 252, 253, 254, 255
54 OTHER INFORMATION: n = A,T,C or G
55 FEATURE:
56 NAME/KEY: misc_feature
57 LOCATION: 256, 257, 258, 259, 260, 261, 262, 263, 264, 265, 266, 267,
58 LOCATION: 268, 269, 270, 271, 272, 273, 274, 275, 276, 277, 278, 279,
59 LOCATION: 280, 281, 282, 283, 284, 285, 286, 287, 288, 289, 290, 291,
60 LOCATION: 292, 293, 294, 295, 296, 297, 298, 299, 300, 301, 302
61 OTHER INFORMATION: n = A,T,C or G
62 US-11-124-368A-2896

```

Db	77735	CACCCCAAGAGTAACCTGCAATTCGA-GTTCAGGAAATTAAGATGCTAAGGCTGTTCT	77793
Qy	121	TGAGAGCCACACAGACGATATATGCTTGCTAAAGTAAGAGGTACTGCTACCTGTG	180
Db	77794	TGAGAGCCACACAGACGATATTTGCTT-CTAAAGTAAGCAGGTACTGCTACCTGTG	77855
Qy	181	GTCTTCACCCCAAGCTGTGTGA-201	
Db	77853	GTCTTCACCCCAAGCTGTGTGA-77873	

Search completed: May 26, 2006, 14:28:23
Job time : 855 secs

Query Match	82.1%	Score 164.6;	DB 15,	Length 91561;
Best Local Similarity	98.0%;	Pred. No. 4.5e-47;		
Matches 197;	Conservative 1;	Mismatches 0;	Indels 3;	Gaps 3;
QY	1	TTGACATCCTCATCTTTCTTAGGTCCTCAAGGCTTCATCTTCAAGGTCATCTGGGTC	60	
Db	77676	TTGACATCCTCATCTTTCTTAGGTCCTC-AGGCTTCATCTTCAAGGTCATCTGGGTC	77734	
QY	61	CACACCAAGAGTGAACCTGCAATCTCATGTTACAGAAATRGAAGTCTTAAGCTTTCT	120	

November 2005

Published_Applications Nucleic Acid and Published_Applications Amino Acid database searches now generate two sets of results each. The Published_Applications databases have been split into two parts to reduce the amount of time required for their daily updates. This results in more machine time being available for processing searches.

Newly published applications will appear in the Published_Applications_New databases; older published applications make up the Published_Applications_Main databases.

Searches run against Nucleic Acid Published_Applications produce two sets of results, with the extensions **.rnpbm** (Published_Applications_NA_Main) and **.rnpbn** (Published_Applications_NA_New):
Searches run against Amino Acid Published_Applications produce two sets of results, with the extensions **.rapbm** (Published_Applications_AA_Main) and **.rapbn** (Published_Applications_AA_New).

OTHER INFORMATION: n = A,T,C or G
US-10-506-549-3

Query Match 13.3%; Score 26.6; DB 6; Length 394191;
Best Local Similarity 71.4%; Pred. No. 50;
Matches 35; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 28 CAAGCCTTATCTCTACAGGTCATCCGGGTCACACCAAGTGA 76
DB 4279 CAAGATCTTCAATTTACAGATCAATTCGGCTGCGAATGAGAAAGGA 4231

RESULT 7

US-10-953-349-15193
Sequence 15193, Application US/10953349
Publication No. US20060107345A1
GENERAL INFORMATION:
APPLICANT: ALEXANDROV, Nikolai et al.
TITLE OF INVENTION: SEQUENCE-DETERMINED DNA FRAGMENTS AND CORRESPONDING POLYPEPTIDES
FILE REFERENCE: 2750-1579PUS2
CURRENT APPLICATION NUMBER: US/10/953,349
CURRENT FILING DATE: 2004-09-30
NUMBER OF SEQ ID NOS: 40252
SOFTWARE: PatentIn version 3.3
SEQ ID NO 15193
LENGTH: 1068
TYPE: DNA
ORGANISM: Glycine max
US-10-953-349-15193

Query Match 13.2%; Score 26.4; DB 6; Length 1068;
Best Local Similarity 47.0%; Pred. No. 5;
Matches 78; Conservative 1; Mismatches 87; Indels 0; Gaps 0;

QY 10 TCATCTTTCTAGGCTCTCAAGGCTTCACTCTCAAGGTATCTGTGGTGACACCAAG 69
DB 186 TTAATGCTCTAGGCTCTCAAGGACACTGGCGGACAGAGTGGCTGGCTCTCA 245
QY 70 AAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTTGGAGCCCA 129
DB 246 CGGTAGACACAACAATATGCGGATGACAGAGTGTCTTCCAGCGCTTCAGAGAGGTGA 305
QY 130 CACACAGATATTTGCTTGAAGCTAAGCAGGTAAGTCTGCTCAC 175
DB 306 AGAAGCCGATCTGSCCATTCATCAATCAATCAATGCTTCTCTCC 351

RESULT 8

US-10-953-349-9738
Sequence 9738, Application US/10953349
Publication No. US20060107345A1
GENERAL INFORMATION:
APPLICANT: ALEXANDROV, Nikolai et al.
TITLE OF INVENTION: SEQUENCE-DETERMINED DNA FRAGMENTS AND CORRESPONDING POLYPEPTIDES
FILE REFERENCE: 2750-1579PUS2
CURRENT APPLICATION NUMBER: US/10/953,349
CURRENT FILING DATE: 2004-09-30
NUMBER OF SEQ ID NOS: 40252
SOFTWARE: PatentIn version 3.3
SEQ ID NO 9738
LENGTH: 3420
TYPE: DNA
ORGANISM: Arabidopsis thaliana
US-10-953-349-9738

Query Match 13.2%; Score 26.4; DB 6; Length 3420;
Best Local Similarity 54.3%; Pred. No. 8.3;
Matches 51; Conservative 1; Mismatches 42; Indels 0; Gaps 0;
QY 51 ATCTGGGTGACACCAAGTGAAGTCTGAATCTCATGTTCAAGAAATGAAAGTGTCT 110

DB 770 ATATGGGTTTCTCGAAGAAATCATCGCCAGTATCAGTTTGGAGATGAAACTTGG 829

QY 111 AGGCTGTTCTTGGAGCCACACAGACAGATATTG 144
DB 830 TAGATGTTTTCAGACCACTTTTCAAGCCGAATTG 863

RESULT 9

US-10-511-937-2815/c
Sequence 2815, Application US/10511937
Publication No. US2006008836A1
GENERAL INFORMATION:
APPLICANT: EXPRESSION DIAGNOSTICS, INC.
APPLICANT: Mohlgemuth, Jay
APPLICANT: Fry, Kirk
APPLICANT: Woodward, Robert
APPLICANT: Ly, Ngoc
APPLICANT: Prentice, James
APPLICANT: Morris, Macdonald
APPLICANT: Rosenberg, Steven
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR DIAGNOSING
TITLE OF INVENTION: AND MONITORING TRANSPLANT REJECTION
FILE REFERENCE: 506612000104
CURRENT APPLICATION NUMBER: US/10/511,937
CURRENT FILING DATE: 2004-10-19
PRIOR APPLICATION NUMBER: PCT/US2003/012946
PRIOR FILING DATE: 2003-04-24
PRIOR APPLICATION NUMBER: US 10/131,831
PRIOR FILING DATE: 2002-04-24
PRIOR APPLICATION NUMBER: US 10/325,899
PRIOR FILING DATE: 2002-12-20
NUMBER OF SEQ ID NOS: 3117
SOFTWARE: PatentIn version 3.2
SEQ ID NO 2815
LENGTH: 6050
TYPE: DNA
ORGANISM: Homo sapiens
US-10-511-937-2815

Query Match 13.2%; Score 26.4; DB 6; Length 6050;
Best Local Similarity 50.0%; Pred. No. 11;
Matches 66; Conservative 0; Mismatches 66; Indels 0; Gaps 0;

QY 1 TTGACATCTCATCTTTTCTAGGCTCTCAAGGCTTATCTCTCAAGTTCATCTGGGTG 60
DB 505 TTGCCATGATCAATTTGAGAGTGAATCTTCTGCTTCAAGTCTTCAAGTCTTGGGT 446
QY 61 CACACCAAGATGAACCTGGAATCTCATGTTCAAGAAATGAAAGTGTCTAGGCTGTCT 120
DB 445 AAGGTTCTGAGAGAGCTGGAGCTTCCATTCATCAATGATGATGATGAGCTGCAC 386
QY 121 TGAAGCCACAC 132
DB 385 TGAAGAAACCC 374

RESULT 10

US-11-293-697-1913/c
Sequence 1913, Application US/11293697
Publication No. US20060105376A1
GENERAL INFORMATION:
APPLICANT: HELIX RESEARCH INSTITUTE
TITLE OF INVENTION: Novel full length cDNA
FILE REFERENCE: H1-A0106
CURRENT APPLICATION NUMBER: US/11/293,697
CURRENT FILING DATE: 2005-12-05
PRIOR APPLICATION NUMBER: US/10/108,260
PRIOR FILING DATE: 2002-03-28
NUMBER OF SEQ ID NOS: 5458
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 1913
LENGTH: 3059
TYPE: DNA

ORGANISM: Homo sapiens
US-11-293-697-1913

Query Match 13.1%; Score 26.2; DB 7; Length 3059;
Best Local Similarity 51.3%; Pred. No. 9.3;
Matches 58; Conservative 1; Mismatches 54; Indels 0; Gaps 0;

QY 15 TTTTCTAGCTCTCAGGCTTTCATCCATCAAGTCACTCGGCGACACCAAGAGT 74
DB 2902 TTTTCTGCTCTCTCTTGTGCAATTAATTAAGAGAAAAGTAATTTAGAGAGAGG 2843

QY 75 AACCTGAACTCATGTTCAAGAAATGAAAGTCTAGGCTGTTCTTGAGGCC 127
DB 2842 GGGCCGCTCTCTCCGATCTGTAATAGTATAGTATGTTGTTCTTGCGCC 2790

RESULT 11
US-11-217-529-79011

Sequence 79011, Application US/11217529
Publication No. US2006009612A1

GENERAL INFORMATION:

APPLICANT: SUNTORY LIMITED

APPLICANT: NAKAO, YOSHIHIRO

APPLICANT: NAKAMURA, NORIHISA

APPLICANT: KODAMA, YUKIKO

APPLICANT: FUJIMURA, TOMOKO

APPLICANT: ASHIKARI, TOSHIHIKO

TITLE OF INVENTION: METHODS FOR ANALYZING GENES OF INDUSTRIAL YEASTS

FILE REFERENCE: S-38-285

CURRENT APPLICATION NUMBER: US/11/217,529

PRIOR FILING DATE: 2005-09-02

PRIOR APPLICATION NUMBER: US 10/932,182

NUMBER OF SEQ ID NOS: 197023

SOFTWARE: PatentIn version 3.3

SEQ ID NO 79011

LENGTH: 1008

TYPE: DNA

ORGANISM: Saccharomyces pastorianus

US-11-217-529-79011

Query Match 12.7%; Score 25.4; DB 7; Length 1008;
Best Local Similarity 53.5%; Pred. No. 11;
Matches 53; Conservative 0; Mismatches 46; Indels 0; Gaps 0;

QY 56 GGGTGCACACCAAGAAAGCTCGAATCTCATGTTCAAGAAATGAAAGTCTAGGCT 115
DB 75 GGATGCAAGAAACGAGTGGAAATGCGAAAAATTCAAAAAGAGCTTCGAATCAATATTC 134

QY 116 GTTCTTGAGGCCACACAGAGATATGCTTGGCTTAA 154
DB 135 GTCAATTAGACGCAAGAGAGCAAAATTTGACTGTATATA 173

RESULT 12

US-11-217-529-76402/C

Sequence 76402, Application US/11217529

Publication No. US2006009612A1

GENERAL INFORMATION:

APPLICANT: SUNTORY LIMITED

APPLICANT: NAKAO, YOSHIHIRO

APPLICANT: NAKAMURA, NORIHISA

APPLICANT: KODAMA, YUKIKO

APPLICANT: FUJIMURA, TOMOKO

APPLICANT: ASHIKARI, TOSHIHIKO

TITLE OF INVENTION: METHODS FOR ANALYZING GENES OF INDUSTRIAL YEASTS

FILE REFERENCE: S-38-285

CURRENT APPLICATION NUMBER: US/11/217,529

PRIOR FILING DATE: 2005-09-02

PRIOR APPLICATION NUMBER: US 10/932,182

NUMBER OF SEQ ID NOS: 197023

SOFTWARE: PatentIn version 3.3

SEQ ID NO 76402

LENGTH: 1734

TYPE: DNA

ORGANISM: Saccharomyces pastorianus

US-11-217-529-76402

Query Match 12.6%; Score 25.2; DB 7; Length 1734;
Best Local Similarity 53.1%; Pred. No. 16;
Matches 51; Conservative 1; Mismatches 44; Indels 0; Gaps 0;

QY 65 CCAAGAGTGAACCTCGAATCTCATGTTCAAGAAATGAAAGTCTAGGCTTCTTGA 124
DB 940 CCAATCTGTAAGCTTCTTATGCTTCTTCAAGACATCATACGTTCAATCTTTTGG 881

QY 125 GCCCACAAGCAGATATGCTTGTCAAGCTAAG 160
DB 880 CACGACCAAGTATACCTTTCTTCATTAGTTGAG 845

RESULT 13

US-11-293-697-740/C

Sequence 740, Application US/11293697

Publication No. US20060105376A1

GENERAL INFORMATION:

APPLICANT: HELIX RESEARCH INSTITUTE

TITLE OF INVENTION: Novel full length cDNA

FILE REFERENCE: H1-A0106

CURRENT APPLICATION NUMBER: US/11/293,697

PRIOR FILING DATE: 2005-12-05

PRIOR APPLICATION NUMBER: US/10/108,260

NUMBER OF SEQ ID NOS: 5458

SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 740

LENGTH: 2638

TYPE: DNA

ORGANISM: Homo sapiens

US-11-293-697-740

Query Match 12.6%; Score 25.2; DB 7; Length 2638;
Best Local Similarity 54.3%; Pred. No. 19;
Matches 51; Conservative 0; Mismatches 43; Indels 0; Gaps 0;

QY 7 TCCATCTTTTCAAGCTCTCAAGGCTTCATCTCAAGTCACTCGGCTGACAGACC 66
DB 1751 TCCCTTTCTGCTCTTTTCCAGTCTGCTGCTGAAGGTGACCTGTACATGA 1692

QY 67 AAGAAGTGAACCTCGAATCTCATGTTCAAGAAAT 100
DB 1691 GATCAGTGAATCTCGGCTCTCAATACCAAGGAGT 1658

RESULT 14

US-10-488-619-2493/C

Sequence 2493, Application US/10488619

Publication No. US2006009578A1

GENERAL INFORMATION:

APPLICANT: Greenlee, Winner and Sullivan, P. C.

TITLE OF INVENTION: Human Mitochondrial DNA Polymorphisms, Haplogroups, Association

FILE REFERENCE: 98-01 WO

CURRENT APPLICATION NUMBER: US/10/488,619

PRIOR FILING DATE: 2004-03-01

NUMBER OF SEQ ID NOS: 3040

SOFTWARE: PatentIn version 3.1

SEQ ID NO 2493

TYPE: DNA

ORGANISM: Mus musculus

US-10-488-619-2493

Query Match 12.5%; Score 25; DB 6; Length 611;
Best Local Similarity 50.4%; Pred. No. 12;

Matches 58; Conservative 1; Mismatches 56; Indels 0; Gaps 0;

QY 81 GAATTCATGTTACAGAAATRGAGTGTCTAGGCTGTTCTTGAGCCCAACAGCAGAT 140

DB 321 GGAGCTGAGTCTGTGAAAGCTGTACAGACTGTTCTCTAGCCCTAACGACGACG 262

QY 141 ATTGCTTGCTTAAGCTAAGCAGTACTGCTCACCTGTGTCTTACCCCAAGC 195

DB 261 CATGATGCTGAGACGACGAGTGTCTCTTCTCTCCGCGGAGAACCCCAAGC 207

RESULT 15

US-10-505-928-226

/ Sequence 226, Application US/10505928

/ Publication No. US20060088532A1

/ GENERAL INFORMATION:

/ APPLICANT: Ludwig Institute for Cancer Research et al.

/ TITLE OF INVENTION: LYMPHATIC ENDOTHELIAL GENES

/ FILE REFERENCE: 28967/39178

/ CURRENT APPLICATION NUMBER: US/10/505,928

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/ SOFTWARE: Patentin 3.2

/ SEQ ID NO: 226

/ LENGTH: 4908

/ TYPE: DNA

/ ORGANISM: Homo sapiens

US-10-505-928-226

Query Match 12.5%; Score 25; DB 6; Length 4908;

Best Local Similarity 59.7%; Pred. No. 29;

Matches 40; Conservative 1; Mismatches 26; Indels 0; Gaps 0;

QY 88 ATGTTACGAAATRGAGTGTCTAGGCTGTTCTTGAGCCCAACAGCAGATATTGCT 147

DB 1079 ATCTAACCTAAGTATGATCTCTCAGGATTTCTTGAGGGCGTACAAACATCTATAAT 1138

QY 148 TGCTAAA 154

DB 1139 TGCAACA 1145

Search completed: May 26, 2006, 14:28:53
Job time : 25 secs